PHYSICAL/CHEMICAL ELEMENTS
1. MELTING POINT

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: No information on source/purity of test material

METHOD

- **Method/guideline:** Japanese MITI test
- **GLP:** No data, but likely to be GLP
- **Year:** 1992

Remarks:

RESULTS

- **Melting point value in °C:** 111-113
- **Decomposition:** No data
- **Sublimation:** No data

Remarks:

CONCLUSIONS

Remarks:

DATA QUALITY

- **Reliabilities:** 2, Valid with restrictions

Remarks: Peer reviewed study, limited information available

REFERENCES (Free Text)

Chemicals and Testing Institute, Biodegradation and Bioaccumulation Data of existing Chemicals, Based on the CSCL Japan, Japan Chemical Industry Ecology-Toxicology and Information Center, ISBN 4-89074-101-1 (1992)

OTHER

- **Last Changed:** October 3 2002
- **Order number for sorting:** 1

Remarks:
-

PHYSICAL/CHEMICAL ELEMENTS
2. BOILING POINT

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)
- Purity: 99.34%
- Lot No.: 023008
- Supplier: Bromine Compounds Ltd.

Remarks:

METHOD

- **Method/guideline:** OPPTS Guidelines Method 830.7220
- **GLP:** Yes
- **Year:** 2003

Remarks:

RESULTS

- **Boiling point value in °C:** 270
- **Pressure:** 100.44 – 101.54 kPa
- **Decomposition:** No

Remarks:

CONCLUSIONS

Remarks: The boiling point of the test material has been determined to be approximately 543K at 100.44 – 101.54 kPa

DATA QUALITY

- **Reliabilities:** 1, Valid without restrictions

Remarks: Guideline study conducted to GLP

REFERENCES (Free Text)

White, D.F., Mullee, D.M., (2003) Safepharm Laboratories Ltd., Dinol: Determination of Boiling Point, OECD Guidelines for Testing of Chemicals, US EPA Office of Prevention, Pesticides and Toxic Substances, Series 830: Product Properties Test Guidelines, Commission Directive 92/69/EEC (which constitutes Annex V of Council Directive 67/548/EEC), Project No. 466/215

OTHER

- **Last Changed:** March, 2004
- **Order number for sorting:** 1

Remarks:

PHYSICAL/CHEMICAL ELEMENTS
3. VAPOUR PRESSURE

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)
- Purity: 99.34%
- Lot No.: 023008
- Supplier: Bromine Compounds Ltd.

METHOD

- Method/guideline: OECD 104 (Vapor pressure balance)
- GLP: Yes
- Year: 2003

RESULTS

- Vapor pressure value: 2.0×10^{-3} Pa
- Temperature: 25°C
- Decomposition: No

CONCLUSIONS

The substance has been determined to have a vapor pressure of 2.0×10^{-3} Pa at 25°C, using a vapor pressure balance

DATA QUALITY

- **Reliabilities:** 1, Valid without restriction

Remarks: Guideline study conducted to GLP

REFERENCES

Tremain, SP (2003), Safepharm Laboratories Ltd., Dinol: Determination of Vapour Pressure, Project No. 466/188

OTHER

- **Last Changed:** March 18, 2004

Order number for sorting: 1

PHYSICAL/CHEMICAL ELEMENTS
4.1 PARTITION COEFFICIENT

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: No information on source/purity of test material

METHOD

- **Method/guideline:** Japanese MITI test
- **GLP:** No data, but likely to be GLP
- **Year:** 1992

Remarks:

RESULTS

- **Log Pow:** 2.29
- **Temperature °C:** Not stated

Remarks:

CONCLUSIONS

Remarks:

DATA QUALITY

- **Reliabilities:** 2, Valid with restrictions

Remarks: Peer reviewed study, limited information available

REFERENCES (Free Text)

Chemicals and Testing Institute, Biodegradation a nd Bioaccumulation Data of existing Chemicals, Based on the CSCL Japan, Japan Chemical Industry Ecology-Toxicology and Information Center, ISBN 4-89074-101-1 (1992)

OTHER

- **Last Changed:** March 26, 2003
- **Order number for sorting:** 1

Remarks:

PHYSICAL/CHEMICAL ELEMENTS
4.2 PARTITION COEFFICIENT

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: No information on source/purity of test material

METHOD

- **Method/guideline:** Hansch, et al
- **GLP:** No data
- **Year:** 1995

Remarks:

RESULTS

- **Log Pow:** 1.06
- **Temperature °C:** Not stated

Remarks: This value is in good agreement with the EPIWIN estimate of 0.85.

CONCLUSIONS

Remarks:

DATA QUALITY

- **Reliabilities:** 2, Valid with restrictions

Remarks: EPA recommended value

REFERENCES (Free Text)

Hansch, C., Leo, A., & Hoekman, D., 1995. Exploring QSAR. Hydrophobic, Electronic and Steric constants. ACS Professional Reference Book. Washington, DC: American Chemical Society

OTHER

- **Last Changed:** March 26, 2003
- **Order number for sorting:** 2

Remarks:

PHYSICAL/CHEMICAL ELEMENTS
5. WATER SOLUBILITY

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)
- Purity: 99.34%
- Lot No.: 023008
- Supplier: Bromine Compounds Ltd.

Remarks:

METHOD

- **Method/guideline:** OPPTS Guidelines, Method 830.7840
- **GLP:** Yes
- **Year:** 2003

Remarks:

RESULTS

- **Value:** 19,400 mg/L at temperature 20.0 ±0.5°C
- **Description of solubility:** Soluble
- **pH value 20°C:** 7.1
- **pKa value at 25°C:** No data

CONCLUSIONS

Remarks: The water solubility of the test material has been determined to be 19.4 g/l of solution at 20.0 ±0.5°C.

DATA QUALITY

- **Reliabilities:** 1, Valid without restrictions

Remarks: Guideline study conducted to GLP

REFERENCES:

White, D.F., Mullee, D.M., (2003) Safepharm Laboratories Ltd., Dinol: Determination of Water Solubility, Data Requirements: OECD Guidelines for Testing of Chemicals, US EPA Office of Prevention, Pesticides and Toxic Substances, Series 830: Product Properties Test Guidelines, Commission directive 92/69/EEC (which constitutes Annex V of Council Directive 67/548/EEC), Project No. 466/216

OTHER

- **Last Changed:** March 18, 2004
- **Order number for sorting:** 1

Remarks:

ENVIRONMENTAL FATE ELEMENTS AND PATHWAYS
6. PHOTODEGRADATION

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

METHOD

- **Method:** Calculated using AOPWIN v1.90
- **GLP:** No
- **Year:** 2003

RESULTS

- **Indirect photolysis**
  - **Sensitizer:** OH radicals
  - **Concentration of sensitizer:** 1.5E+06 OH/cm^3
  - **Rate constant:** 8.9651E-12 cm^3/molecule-sec
  - **Half-life:** 14.3 hours

CONCLUSIONS

DATA QUALITY

- **Reliabilities:** 1, valid without restrictions

REFERENCES (Free Text)

AOPWIN v1.90. EPIWIN Modelling Program. Meylan, W. & Howard, P. (2000), Syracuse Research Corporation, Environmental Science Center, 6225 Running Ridge Road, North Syracuse, NY 13212-2510

OTHER

- January 5, 2004
- 1

ENVIRONMENTAL FATE ELEMENTS AND PATHWAYS
7. STABILITY IN WATER

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)
- Purity: 99.34%
- Lot No.: 023008
- Supplier: Bromine Compounds Ltd.

METHOD

- Method/guideline: OECD 111
- GLP: Yes
- Year: 2003

Remarks:

- **Duration:** 120 – 144 hours @ 50°C
- **Analytical procedures:** The standard and sample solutions were analysed by HPLC using the following conditions:

HPLC System: Agilent Technologies 1100 or 1050, incorporating autosampler and workstation
Column: Prodigy 5µ ODS (2) (250 x 4.6 mm i.d.)
Column temperature: 40°C
Mobile phase: methanol: reverse osmosis water (55:45 v/v)
Flow-rate: 1.0 ml/min
UV detector wavelength: 210 nm
Injection volume: 20 µl
Retention time: ~ 9 mins

RESULTS

- **Nominal concentration:** 1 g/l
- **Measured value:** 1.01 – 1.04 g/l
- **Half-life:** See table below

pH		Time at 50°C (hours)					% hydrolysis at 50°C	Half-life at 25°C
		0	2.4	24	120	144		
4	Concentration (g/l)	1.01	1.01	0.892	0.873	1.02		
	% of initial	-	101	88.5	86.6	102	< 10%	> 1yr
7	Concentration (g/l)	1.04	1.06	1.03	1.02	-		
	% of initial	-	102	99.8	98.8	-	< 10%	> 1yr
9	Concentration (g/l)	1.04	1.04	1.01	0.897	-		
	% of initial	103	103	99.8	88.9	-	~ 10%	~ 1 yr

- **Breakdown products:** No

Remarks: An extra timepoint was taken for pH 4 at 144 hours. This was due to the unexpectedly low value obtained for the 120 hour sample.

CONCLUSIONS

The estimated half-life of the substance at 25° is > 1 year at pH 4 and 7 and approximately equal to 1 year at pH 9.

DATA QUALITY

- **Reliabilities:** 1, Valid without restriction

Remarks: Guideline study conducted to GLP

REFERENCES

White, DF & Mullee, DM (2003), Safepharm Laboratories Ltd., Dinol: Determination of Hydrolysis as a function of pH, Project No. 466/255

OTHER

- **Last Changed:** September 22, 2003
- **Order number for sorting:** 1

ENVIRONMENTAL FATE ELEMENTS AND PATHWAYS
8. TRANSPORT BETWEEN ENVIRONMENTAL COMPARTMENTS (FUGACITY)

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

METHOD

- **Test:** Level III Fugacity Model
- **Method:** Calculated using Epiwin Level III Fugacity Model
- **Year:** 2003
- **GLP:** No

Remarks

Henry's Law constant: 2.66E-10 atm-m^3/mole (calc VP/Wsol)
Vapor Pressure: 1.5E-05 mm Hg (user entered)
Liquid VP: 0.000106 mm Hg (super cooled)
Melting Point: 111°C (user entered)
Log Kow: 1.06 (Kowwin program)
Soil Koc: 4.71 (calc by model)

RESULTS

	Mass Amount (percent)	Half-life (hr)	Emissions (kg/hr)
Air	0.00855	28.6	1000
Water	41.4	360	1000
Soil	58.5	360	1000
Sediment	0.0759	1440	0

	Fugacity (atm)	Reaction (kg/hr)	Advection (kg/hr)	Reaction (percent)	Advection (percent)
Air	1.01E-13	2.65	1.1	0.0884	0.0365
Water	2.7E-15	1020	531	34.1	17.7
Soil	1.03E-13	1440	0	48.1	0
Sediment	2.22E-15	0.468	0.0194	0.0156	0.000648

Persistence time: 427 hr
Reaction time: 519 hr
Advection time: 2410 hr
Percent reacted: 82.3
Percent advected: 17.7

Half-lives (hr), (based upon Biowin (ultimate) and Aopwin):
Air: 28.63
Water: 360
Soil: 360
Sediment: 1440

Biowin estimate: 2.786 (months)

Advection times (hr):
Air: 100
Water: 1000
Sediment: 5E+4

CONCLUSIONS

DATA QUALITY

- **Reliabilities:** 1, valid without restrictions

REFERENCES (Free Text)

Level III Fugacity Model. EPIWIN Modelling Program. Meylan, W. & Howard, P. (2000), Syracuse Research Corporation, Environmental Science Center, 6225 Running Ridge Road, North Syracuse, NY 13212-2510

OTHER

- **Last changed:** January 05, 2004
- **Order number for sorting:** 1

ENVIRONMENTAL FATE ELEMENTS AND PATHWAYS
9. BIODEGRADATION

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: No information on source/purity of test material

METHOD

- **Method/guideline:** Japanese MITI test
- **Test Type:** Aerobic
- **GLP:** No data, but likely to be GLP
- **Year:** 1992
- **Contact time (units):** 28 days
- **Innoculum:** Activated sewage sludge

Remarks:

- **Innoculum (concentration and source)**
  - **Fresh activated sludge:** 30 mg/l

- **Concentration of test chemical:** 100 mg/l
- **Temperature of incubation °C:** No data
- **Dosing procedure:** No data
- **Sampling frequency:** No data
- **Appropriate controls and blank system used?:** No data
- **Analytical method used to measure biodegradation:** No data
- **Method of calculating measured concentrations:** No data

RESULTS

- **Degradation % after time:** 3 – 33% by BOD after 28 days
- **Results:** Not readily biodegradable
- **Kinetic:** No data

Breakdown products (yes/no): No data

Remarks:

CONCLUSIONS

Remarks: Based on this test, the substance is not readily biodegradable

DATA QUALITY

- **Reliabilities:** 2, Valid with restrictions

Remarks: Peer reviewed study, limited information available

REFERENCES (Free Text)

Chemicals and Testing Institute, Biodegradation and Bioaccumulation Data of existing Chemicals, Based on the CSCL Japan, Japan Chemical Industry Ecology-Toxicology and Information Center, ISBN 4-89074-101-1 (1992)

OTHER

- **Last Changed:** October 3 2002
- **Order number for sorting:** 1

Remarks:

ECOTOXICITY ELEMENTS
10. TOXICITY TO FISH

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)
- Purity: 99.34%
- Lot No.: 023008
- Supplier: Bromine Compounds Ltd.

Remarks:

METHOD

- **Method/guideline:** OECD 203
- **Type:** Semi-static
- **GLP:** Yes
- **Year:** 2003
- **Species and source:** Oncorhynchus mykiss. Brow Well Fisheries Ltd., Hebden, Yorkshire, UK
- **Analytical monitoring:** Yes
- **Exposure period:** 96 hours
- **Statistical methods:** none

Remarks:

Test fish: The test was performed using juvenile fish. At the end of the test the fish had a mean standard length of 4.9±0.4 cm, and a mean weight of 1.52±0.43 g. Based on the mean weight value, the loading rate was 0.38 g bodyweight/litre.

Test Conditions:

- **Dilution water source:** Laboratory tap water was dechlorinated by passage through as activated carbon filter (Purite Series 500) and partly softened (Elga Nimbus 1248D duplex water softener).
- **Dilution water chemistry:** Hardness = 100 mg/L as $CaCO_3$, pH = 7.5-7.6
- **Stock and test solution preparation:** An amount of test material (8.00 g) was dissolved in dechlorinated tap water with the aid of ultrasonication (approximately 15 minutes) and the volume adjusted to 2 litres to give a 4 g/L stock solution. An aliquot (1 litre) of the stock solution was dispersed in dechlorinated tap water and the volume adjusted to 4 0 litres to give the 100 mg/L test concentration.
- **Concentrations:** 100 mg/L in the definitive test
- **Vehicle/solvent:** None
- **Stability of the test chemical solutions:** Preliminary test samples were prepared, analysed initially and then after storage in sealed glass vessels at ambient temperature in light and dark conditions for approximately 24 hours (equivalent to the period of media renewal). In addition a test sample was tested for stability without prior mixing (sonication) of the test sample bottle to assess for losses due to absorption and/or insolubility. The test samples were shown to be stable in the test medium and the unsonicated stability vessel showed no evidence of insolubility or adherence to glass.
- **Exposure vessel type:** 40 litre glass exposure vessels were used for each test concentration. The test vessels were covered to reduce evaporation. A photoperiod of 16 hours light and 8 hours darkness with 20 minute dawn and dusk transition periods was used. The test vessels were aerated via narrow bore glass tubes.
- **Number of replicates, fish per replicate:** 2 replicates per test concentration, 10 fish per replicate

- **Water chemistry in test, test temperature range:** See table below

Nominal concentration (mg/L)	Maximum & minimum values measured over the 96 hour test period		
	pH	mg O_2/L	T°C
Control	7.5 – 7.9	8.8 – 9.3	14.6 – 14.8
100 (Replicate 1)	7.5 – 8.1	8.9 – 9.6	14.6 – 14.9
100 (Replicate 2)	7.5 – 8.1	8.8 – 9.7	14.6 – 14.8

RESULTS

- **Nominal/measured concentrations (as mg/L):** See table below

Sample time (hours)	Nominal concentration (mg/L)	Measured concentration (mg/L)	Percent of nominal (%)
0	Control	<LOQ	-
	100 (Replicate 1)	103	103
	100 (Replicate 2)	110	110
24	Control	<LOQ	-
	100 (Replicate 1)	101	101
	100 (Replicate 2)	99.0	99
96	Control	< LOQ	-
	100 (Replicate 1)	104	104
	100 (Replicate 2)	103	103

Analysis of the test preparations at 0, 24 and 96 hours showed measured test concentrations to range from 99 – 110% of nominal and so results were based on nominal test concentrations only.

- **Unit:** mg/L
- **Element value:** LC_{50}
- **Result:** LC_{50} > 100 mg/L

Remarks:

Biological observations: There were no sub-lethal effects of exposure in 20 fish exposed to a test concentration of 100 mg/L for a period of 96 hours.

Cumulative mortality: See table below.

Nominal Concentration (mg/L)	Cumulative mortality						% Mortality
	3 hours	6 hours	24 hours	48 hours	72 hours	96 hours	96 hours
Control	0	0	0	0	0	0	0
100 (Replicate 1)	0	0	0	0	0	0	0
100 (Replicate 2)	0	0	0	0	0	0	0

CONCLUSIONS

Remarks: The results of this study indicate that, at concentrations up to and including 100 mg/l, the maximum concentration given in the OECD guideline, the test material was not toxic to fish.

DATA QUALITY

- **Reliabilities:** 1, Valid without restriction

Remarks: Guideline study conducted to GLP

REFERENCES (Free Text)

Sewell, I.G. & McKenzie, J (2003), Safepharm Laboratories Ltd., Dinol: Acute toxicity to Rainbow Trout (Oncorhynchus mykiss), Project No. 466/191

OTHER

- **Last Changed:** March 18, 2004
- **Order number for sorting:** 1

Remarks:

ECOTOXICITY ELEMENTS
11. TOXICITY TO AQUATIC PLANTS (E.G., ALGAE)

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)
- Purity: 99.34%
- Lot No.: 023008
- Supplier: Bromine Compounds Ltd.

Remarks:

METHOD

- **Method/guideline:** OECD 201
- **Type:** Static
- **GLP:** Yes
- **Year:** 2003
- **Species/strain # and source:** Scenedesmus subspicatus CCAP 276/20, Culture Collection of Algae and Protozoa (CCAP), Institute of Freshwater Ecology, United Kingdom.
- **Element basis:** Growth rate, biomass
- **Exposure period:** 72 hours
- **Analytical monitoring:** Yes
- **Statistical methods:** One way analysis of variance incorporating Bartlett's test for homogeneity of variance (Sokal & Rohlf 1981) and Dunnett's multiple comparison procedure for comparing several treatments with a control (Dunnett 1955) was carried out on the area under the growth curve data at 72 hours for the control and all test concentrations to determine any statistically significant differences between the test and control groups. All statistical analyses were performed using the SAS computer software package (SAS 1999 – 2001).

Dunnett, CW (1955) A multiple comparison procedure for comparing several treatments with a control. J Am Stat Assoc, 50, 1096-1121

SAS/STAT proprietary software release 8.02 (1999-2001), SAS Institute Inc., Cary, NC, USA

Sokal, RR and Rohlf, FJ (1981) Biometry. New York: WH Freeman and Company

Remarks:

Test organisms: Liquid cultures of Scenedesmus subspicatus were obtained from CCAP. Cultures were maintained in the laboratory by the periodic replenishment of culture medium. The culture was maintained in the laboratory at a temperature of 21±1°C under continuous illumination (intensity approximately 7000 lux) and constant aeration.

Test Conditions:

- **Test temperature range:** 23 – 25°C
- **Dilution water source:** Reverse osmosis purified deionised water (Elga Optima 15+)
- **Exposure vessel type:** 250 ml glass conical flasks (3 per group) each containing 100 ml of test preparation, plugged with polyurethane foam bungs.
- **Water chemistry in test (pH):** See table below

Nominal concentration (mg/L)		pH	
		0 Hours	72 Hours
Control	R1	7.4	8.4
	R2	7.4	8.4
	R3	7.4	8.4
6.25	R1	7.4	8.3
	R2	7.4	8.3
	R3	7.4	8.3
12.5	R1	7.3	8.2
	R2	7.3	8.2
	R3	7.3	8.2
25	R1	7.2	8.3
	R2	7.2	8.3
	R3	7.2	8.3
50	R1	7.2	8.2
	R2	7.2	8.2
	R3	7.2	8.2
100	R1	7.1	8.1
	R2	7.1	8.1
	R3	7.1	8.1

- **Stock solutions preparation:** An amount of test material (100 mg) was dissolved in culture medium with the aid of ultrasonic disruption (approximately 10 minutes) and the volume adjusted to 500 ml to give a 200 mg/l stock solution from which serial dilutions were made to prepare 100, 50, 25 and 12.5 mg/l stock solutions. An aliquot (250 ml) of each stock solution was separately mixed with algal suspension (250 ml) to give the required test concentrations of 6.25, 12.5, 25, 50 and 100 mg/l.
- **Light levels and quality during exposure:** 7000 lux, continuous illumination

Test design (number of replicates, concentrations): 3 replicates per test concentration. 0, 6.25, 12.5, 25, 50, 100 mg/L

RESULTS

- **Nominal/measured concentrations (as mg/L):** See table below

Nominal Concentration (mg/L)	Sample Time			
	0 Hours		72 Hours	
	Measured Concentration (mg/L)	Percent of nominal (%)	Measured Concentration (mg/L)	Percent of nominal (%)
0	<LOQ	-	<LOQ	-
6.25	4.81	77*	5.98	96
12.5	10.9	87	13.0	104
25	22.3	89	25.0	100
50	50.0	100	51.7	103
100	93.4	93	92.6	93

*This low result was considered to be due to sampling/analytical variation as all other results were in the accepted range of 80-100%
LOQ = 0.31 mg/l

Analysis of the test solutions at 0 and 72 hours showed measured test concentrations to be near nominal with the exception of the 6.25 mg/l test group at 0 hours which gave a measured concentration of 77% of nominal. This low measured concentration was considered to be due to sampling and/or analytical variation as all other measured test concentrations at 0 and 72 hours were near nominal. It was therefore considered appropriate to base the results on nominal concentrations.

- **E_bC_{50}(72h) = 37 mg/L, E_rC_{50}(0-72h) = 150 mg/L**

E_rC_{50} value determined from the equation for the fitted line, as no concentration tested resulted in 50% inhibition of growth rate.

- **NOEC = 12.5 mg/L**

- **Statistical results, as appropriate:** Statistical analysis of the area under the growth curve data was carried out for the control and all test concentrations. There were no statistically significant differences between the control and 6.25 and 12.5 mg/l test concentrations ($p \geq 0.05$), however all other test concentrations were significantly different ($p < 0.05$).

Remarks:

Biological observations: All test and control cultures were inspected microscopically at 72 hours. There were no abnormalities detected in any of the control or test cultures at 6.25, 12.5 and 25 mg/l, however some clumping of algal cells was observed in the test cultures at 50 and 100 mg/l.

- **Cell density at each flask at each measuring point:** See table below

Nominal Concentration (mg/L)		Cell Densities* (cells per mL)			
		0h	24h	48h	72h
Control	R1	9.30E+03	7.84E+04	6.42E+05	1.13E+06
	R2	9.10E+03	7.34E+04	6.58E+05	1.15E+06
	R3	8.76E+03	7.34E+04	6.50E+05	1.10E+06
	Mean	9.05E+03	7.51E+04	6.50E+05	1.13E+06
6.25	R1	9.44E+03	7.67E+04	6.17E+05	1.17E+06
	R2	8.56E+03	7.84E+04	6.33E+05	1.13E+06
	R3	1.09E+04	8.17E+04	6.58E+05	1.13E+06
	Mean	9.64E+03	7.89E+04	6.36E+05	1.14E+06
12.5	R1	1.02E+04	7.67E+04	6.33E+05	1.00E+06
	R2	9.30E+03	7.84E+04	6.33E+05	1.00E+06
	R3	9.56E+03	7.34E+04	6.83E+05	1.07E+06
	Mean	9.70E+03	7.62E+04	6.50E+05	1.02E+06
25	R1	9.78E+03	6.00E+04	5.00E+05	1.05E+06
	R2	9.97E+03	5.84E+04	4.67E+05	1.05E+06
	R3	9.50E+03	6.00E+04	4.83E+05	1.05E+06
	Mean	9.75E+03	5.95E+04	4.83E+05	1.05E+06
50	R1	9.19E+03	4.34E+04	1.83E+05	3.17E+05
	R2	9.43E+03	3.67E+04	1.63E+05	3.00E+05
	R3	9.34E+03	4.34E+04	1.83E+05	3.17E+05
	Mean	9.32E+03	4.12E+04	1.76E+05	3.11E+05
100	R1	9.12E+03	2.17E+04	8.84E+04	2.23E+05
	R2	8.82E+03	1.84E+04	8.84E+04	2.23E+05
	R3	8.97E+03	2.17E+04	8.34E+04	2.13E+05
	Mean	8.97E+03	2.06E+04	8.67E+04	2.20E+05

Cell densities represent the mean number of cells per ml calculated from the mean of cell counts from 3 counts or fields of view for each of the replicate flasks.

- **Percent biomass/growth rate inhibition per concentration:** See table below

Nominal Concentration (mg/L)	Area Under Curve at 72h	% Inhibition	Growth Rate (0-72 h)	% Inhibition
0	3.04E+07	-	0.067	-
6.25	3.03E+07	0	0.066	1
12.5	2.91E+07	4	0.065	3
25	2.50E+07	18	0.065	3
50	8.40E+06	72	0.049	27
100	4.67E+06	85	0.044	34

CONCLUSIONS

Remarks: The effect of test material on the growth of algae has been investigated and gave an E_bC_{50} (72h) value of 37 mg/l and an ErC50 (0-72h) value of 150 mg/l and No Observed Effect Concentration value of 12.5 mg/l.

DATA QUALITY

- **Reliabilities:** 1, Valid without restriction

Remarks: Guideline study conducted to GLP

REFERENCES (Free Text)

Mead, C. & McKenzie, J (2003), Safepharm Laboratories Ltd., Dinol: Algal Inhibition Test, Project No. 466/193

OTHER

- **Last Changed:** March 18, 2004
- **Order number for sorting:** 1

Remarks:

ECOTOXICITY ELEMENTS
12. TOXICITY TO AQUATIC INVERTIBRATES (E.G., DAPHNIA)

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)
- Purity: 99.34%
- Lot No.: 023008
- Supplier: Bromine Compounds Ltd.

METHOD

- **Method/guideline:** OECD 202
- **Type:** Static
- **GLP:** Yes
- **Year:** 2003
- **Analytical procedures:** Water samples were taken at 0 and 48 hours for quantitative analysis by HPLC using an external standard (see below for details).
- **Species/strain:** Daphnia magna
- **Statistical methods:** Not applicable

Remarks:

Test organisms

- **Source:** 1st instar daphnia magna derived from in-house laboratory cultures
- **Age at study initiation:** < 24 hours old
- **Control group:** yes

Test Conditions

- **Stock solutions preparation:** 200 mg of test material was dissolved in reconstituted water with the aid of ultrasonication for approximately 5 minutes, prior to adjusting the volume to 2 liters to give the 100 mg/L test concentration. Aliquots of the 100 mg/L test concentration were each separately dispersed in reconstituted water and the volume adjusted to 1 liter to give the individual test concentrations.
- **Test temperature range:** 20.7 – 21.7°C
- **Exposure vessel type:** 250 mL glass jars containing approximately 200 mL of test preparation, covered to reduce evaporation.
- **Dilution water source:** Reconstituted water
- **Dilution water chemistry:** Total hardness = 250 mg/L as CaCO3; pH 7.8±0.2; oxygen approximately at air saturation value
- **Lighting:** 16 hours light, 8 hours darkness with 20-minute dawn and dusk transition periods.
- **Water chemistry in test:** pH 7.8 – 8.0; O_2 saturation 8.2 – 8.5 mg O_2/L
- **Element (unit) basis:** Immobilization
- **Test design:** Duplicate tests; 10 daphnids per replicate; 0, 1.0, 1.8, 3.2, 5.6, 10, 18, 32, 56 and 100 mg/L.
- **Method of calculating mean measured concentrations:** Not applicable
- **Exposure period:** 48 hours
- **Analytical monitoring:** HPLC, conditions below

HPLC System:
Agilent Technologies 1100 LC/MSD, incorporating autosampler and workstation
Mass selective detector
Source: electrospray
Fragmentation energy: 60 volts
Polarity: negative
Mode: single ion mode with 307 amu
Gas temperature: 250°C
Drying gas: 12 L/min
Nebuliser pressure: 50 psi
Capillary voltage: 2000 volts
Gain: 1
Column: Luna C18 (2), 3µ (100 x 2 mm id)
Column temperature: 40°C
Mobile phase: water:acetonitrile (80:20 v/v)
Flow rate: 0.5 mL/min
Injection volume: 50 µL
Retention time: approximately 3 minutes

RESULTS

- Nominal concentrations: See table below
- Measured concentrations: See table below

Nominal and Measured concentrations found during the study:

Nominal concentration (mg/L)	Measured concentration at 0 hours (mg/L)	Percentage of Nominal (%)	Measured concentration at 48 hours (mg/L)	Percentage of Nominal (%)
Control	<LOQ	-	<LOQ	-
1.0	1.16	116	1.02	102
1.8	N.M.	-	N.M.	-
3.2	3.51	110	3.25	102
5.6	N.M	-	N.M	-
10	10.5	105	9.78	98
18	N.M.	-	N.M.	-
32	32.8	103	31.0	97
56	N.M.	-	N.M.	-
100	106	106	94.0	94

N.M.: Not measured
LOQ: Limit of Quantitation

- **Unit:** mg/L
- EC_{50} > 100 mg/L
  NOEC = 56 mg/L

Remarks:

Biological observations

- **Cumulative immobilization:** See table below
- **Was control response satisfactory:** Yes

Nominal Concentration (mg/L)	Cumulative Immobilzed Daphnia (Initial Population: 10 per Replicate)							
	24 hours				48 hours			
	R_1	R_2	Total	%	R_1	R_2	Total	%
Control	0	0	0	0	0	0	0	0
1.0	0	0	0	0	0	0	0	0
1.8	0	0	0	0	0	0	0	0
3.2	0	0	0	0	0	0	0	0
5.6	0	0	0	0	0	0	0	0
10	0	0	0	0	0	0	0	0
18	0	0	0	0	0	0	0	0
32	0	0	0	0	0	0	0	0
56	0	0	0	0	0	0	0	0
100	2	3	5	25	3	4	7	35

CONCLUSIONS

The acute toxicity of the test material to the freshwater invertebrate Daphnia magna has been investigated and gave a 48 hour EC_{50} value of greater than 100 mg/L. The No Observed Effect Concentration at 48 hours was 56 mg/L.

DATA QUALITY

- **Reliabilities:** 1, Valid without restriction

Remarks: Guideline study conducted to GLP

REFERENCES (Free Text)

Sewell, I.G.& McKenzie, J (2003), Safepharm Laboratories Ltd., Dinol: Acute Toxicity to Daphnia Magna, Project No. 466/192

OTHER

- **Last Changed:** September 26, 2003
- **Order number for sorting:** 1

HUMAN HEALTH ENDPOINTS
13.1 ACUTE TOXICITY

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Bromine Compounds Ltd.
Lot No.: 3456, Purity: 98.96%

METHOD

- **Method:** OECD 401
- **Test Type:** OECD single dose oral toxicity (limit) test
- **GLP:** Yes
- **Year:** 1992
- **Species/strain:** Rat, Sprague-Dawley
- **Sex:** Male/female
- **No. of animals/sex/dose:** Range finding study: 1 male, 1 female
  Main study: 5 males, 5 females
- **Vehicle:** Arachis oil
- **Route of administration:** Oral (single dose gavage)

Remarks:

- **Age:** 5 – 8 weeks old (bodyweight 136 – 153g males, 136 – 146g females)
- **Doses:** 2000 mg/kg
- **Volume administered:** 10 ml/kg
- **Post dose observation period:** Range finding study: 5 days
  Main study: 14 days

RESULTS

- **LD50:** > 2000 mg/kg bodyweight
- **Number of deaths at each dose level:**
  Range finding study: No deaths
  Main study: See Table 1. Three animals (one male and two females) were found dead after dosing at 2000 mg/kg.

Remarks:

- **Time of death:** Main study: Two females died 30 minutes after dosing. One male died one hour after dosing.
- **Clinical signs at each dose level:**
  Range finding study: Common signs of systemic toxicity noted were hunched posture, lethargy, ataxia, laboured respiration, decreased respiratory rate and ptosis with additional signs of pilo-erection, loss of righting reflex, red/brown staining around the eyes, vocalisation and increased lachrymation.

  Main study: See Table 1. Common signs of systemic toxicity noted were coma, laboured respiration, hunched posture and lethargy with additional signs of ataxia, ptosis and decreased respiratory rate.

Isolated incidents of loss of righting reflex were noted in two animals. Surviving animals appeared normal one or two days after dosing.

Table 1: Individual clinical observations and mortality data

Dose Level mg/kg	Animal number & sex	Effect noted after dosing (hours)				Effects noted after dosing (days)													
		1/2	1	2	4	1	2	3	4	5	6	7	8	9	10	11	12	13	14
2000	3-0 male	Co,Rl	X																
	3-1 male	H,L,A,Rl	H,L,A,Rl	H,L,A.Rl	H,L	0	0	0	0	0	0	0	0	0	0	0	0	0	0
	3-2 male	Co,Rl	Co,Pt Rl,Rd	Rr,Rl,Rd	H,L	0	0	0	0	0	0	0	0	0	0	0	0	0	0
	3-3 male	H,L,A,Rl	H,L,A Rl,Pt	H,L,A Rl,Pt	H,L	0	0	0	0	0	0	0	0	0	0	0	0	0	0
	3-4 male	Co,Rl	Co,Rl,Rd	Co,Rl,Rd	Co,Pt Rd,Pt	H,L	0	0	0	0	0	0	0	0	0	0	0	0	0
	4-0 female	X																	
	4-1 female	Co,Rl	Co,Rl,Rd	Co,Rl Rd,Pt	Co,Pt Rd,Rl	H,L	0	0	0	0	0	0	0	0	0	0	0	0	0
	4-2 female	H,L,A	H,L,A,Rl	H,L,A,Rl	H,L,A	L	0	0	0	0	0	0	0	0	0	0	0	0	0
	4-3 female	Rr,L,Rl	Rr,Rl,Rd	H,L,A,Rl	H,L,Rl,A	L	0	0	0	0	0	0	0	0	0	0	0	0	0
	4-4 female	X																	

A – ataxia, Co – coma, H – hunched posture, L – lethargy, Pt – ptosis, Rd – decreased respiratory rate, Rl – laboured respiration, Rr – loss of righting reflex, 0 – no signs of systemic toxicity, X – animal dead

- **Necropsy findings:** See Table 2. Abnormalities noted at necropsy of animals that died during the study were dark liver, dark kidneys and slight haemorrhage of the gastric mucosa. No abnormalities were noted at necropsy of animals that were killed at the end of the study.

Table 2: Individual necropsy findings

Dose Level mg/kg	Animal Number & Sex	Time of Death	Macroscopic Observations
2000	3-0 male	Found dead day 0	Liver: dark Kidneys: dark Gastric mucosa: slight haemorrhage
	3-1 male	Killed day 14	No abnormalities detected
	3-2 male	Killed day 14	No abnormalities detected
	3-3 male	Killed day 14	No abnormalities detected
	3-4 male	Killed day 14	No abnormalities detected
	4-0 female	Found dead day 0	Liver: dark Kidneys: dark Gastric mucosa: slight haemorrhage
	4-1 female	Killed day 14	No abnormalities detected
	4-2 female	Killed day 14	No abnormalities detected
	4-3 female	Killed day 14	No abnormalities detected
	4-4 female	Found dead day 0	Liver: dark Kidneys: dark

CONCLUSIONS

Remarks: The acute median lethal dose (LD_{50}) was found to be greater than 2000 mg/kg bodyweight.

DATA QUALITY

- **Reliabilities:** 1, Reliable without restriction

Remarks: Study conducted under GLP to OECD test guideline by SafePharm Laboratories Ltd.

REFERENCES (Free Text)

SafePharm Laboratories Ltd., DBNPG (Dibromoneopentyl glycol): Acute oral toxicity (Limit test) in the rat, Report No. 466/3, June 1992

OTHER

- **Last Changed:** March 2004
- **Order number for sorting:** 1

Remarks:

HUMAN HEALTH ENDPOINTS
13.2 ACUTE TOXICITY

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Dead Sea Bromine Company Ltd

METHOD

- **Method/guideline followed:** Similar to OECD 401
- **Type:** Oral gavage LD_{50} study in rat
- **GLP:** No, but appears to be conducted to GLP principles
- **Year:** 1979
- **Species/strain:** Rat, CD strain
- **Sex:** Male/female
- **No of animals/sex/dose:** 5 male, 5 female per dose group
- **Vehicle:** Distilled water
- **Route of administration:** Oral (single dose gavage)

Remarks:

- **Age:** Approximately 35 days old (bodyweight 111 – 144g males, 106 – 129g females)
- **Doses:** 1247, 1484, 1765, 2101, 2500 mg/kg
- **Volume administered or concentration:** 20 ml/kg
- **Post dose observation period:** 14 days

RESULTS

- **LD50:** 1880 mg/kg, 95% confidence limits 1691 – 2120 mg/kg
- **Number of deaths at each dose level:** See Table 1

Table 1: Deaths at each dose level

Dosage (mg/kg)	Mortality		
	Male	Female	Combined
1247	0/5	0/5	0/10
1484	0/5	1/5	1/10
1765	2/5	4/5	6/10
2101	3/5	4/5	7/10
2500	4/5	4/5	8/10

Remarks:

- **Time of death:** See Tables 2 - 6
- **Clinical signs at each dose level:** See Tables 2 – 6. Initials signs of reaction to treatment consisted of a rapid progression from reduced motor activity, ataxia, prone posture and prostration to a loss of consciousness between 15 and 30 minutes after administration. Coma and death, between one and six hours after dosing, followed loss of consciousness in a total of 22 animals. The remaining animals generally regained consciousness one to two hours after administration of the lowest dosage, but recovery was less rapid as the dosage increased. All surviving animals were

asymptomatic from Day 3 and all showed normal bodyweight gains over the 14-day observation period.

Table 2: Time of death and clinical signs at 1247 mg/kg dose

Dosage mg/kg	Signs of reaction to treatment	Animal number and sex										Number showing effect at time after dosing														
		A850 Male					A851 Female					Hour						Day								
		1	2	3	4	5	1	2	3	4	5	1/4	1/2	1	2	4	6	2		3	4	5	6	7	8-15
																		am	pm						
1247	**Early decedents** (none)																								
	No. of animals											0	0	0	0	0	0	0	0	0	0	0	0	0	0
	Animals surviving to Day 15:																								
	None															4	4	10	10	10	10	10	10	10	10
	Piloerection					+		+	+		+			1	1	2	3								
	Abdominal hunching			+	+					+				2	2	1									
	Hypoactivity		+		+	+	+	+	+	+	+				3	3	5								
	Inactivity		+	+	+	+				+		4	3	4	2										
	Ataxia		+	+	+	+				+		4	3	4	1										
	Unconsciousness	+					+	+	+	+	+		5	5	3	2	1								
	Proneness		+	+	+	+						4	3	1											
	Prostration	+	+				+	+	+	+	+	6	2	1	2	1									
	No. of animals											10	10	10	10	10	10	10	10	10	10	10	10	10	10

+ Indicates animal exhibiting sign

Table 3: Time of death and clinical signs at 1484 mg/kg dose

Dosage mg/kg	Signs of reaction to treatment	Animal number and sex										Number showing effect at time after dosing														
		A852 Male					A853 Female					Hour						Day								
		1	2	3	4	5	1	2	3	4	5	1/4	1/2	1	2	4	6	2		3	4	5	6	7	8-15
																		am	pm						
1484	**Early decedents**																								
	Unconsciousness							+				1	1	1											
	Death							+							1	1	1	1	1	1	1	1	1	1	1
	Time of death (hours)							2																	
	No. of animals											1	1	1	1	1	1	1	1	1	1	1	1	1	1
	Animals surviving to Day 15:																								
	None																	9	9	9	9	9	9	9	9
	Piloerection			+		+	+		+		+				1	4	5								
	Abdominal hunching		+										1			1	1								
	Hypoactivity		+	+		+	+		+		+	1			2	3	5								
	Inactivity		+	+		+							1	1	1	2									
	Ataxia		+				+		+		+	1	1	1	2		1								
	Loss of righting reflex			+									1												
	Unconsciousness	+			+	+	+		+	+	+	4	7	6	5	4	3								
	Prostration	+		+	+	+			+			4	1	2	1										
	Proneness								+						1										
	No. of animals											9	9	9	9	9	9	9	9	9	9	9	9	9	9

+ Indicates animal exhibiting sign

Table 4: Time of death and clinical signs at 1764 mg/kg dose

Dosage mg/kg	Signs of reaction to treatment	Animal number and sex										Number showing effect at time after dosing														
		A854 Male					A855 Female					Hour						Day								
		1	2	3	4	5	1	2	3	4	5	1/4	1/2	1	2	4	6	2		3	4	5	6	7	8-15
																		am	pm						
1765	**Early decedents**																								
	Unconsciousness	+				+	+	+	+		+	6	6	2	1	1									
	Death	+				+	+	+	+		+			4	5	5	6	6	6	6	6	6	6	6	6
	Time of death (hours)	1				2	6	1	1		1														
	No. of animals											6	6	6	6	6	6	6	6	6	6	6	6	6	6
	Animals surviving to Day 15:																								
	None														1	1	1	4	4	4	4	4	4	4	4
	Hypoactivity			+										1											
	Inactivity			+								1	1												
	Ataxia			+								1	1	1											
	Unconsciousness		+		+					+		3	3	3	3	3	2								
	Proneness			+								1	1				1								
	Prostration				+																				
	Hypopnoea		+											1											
	No. of animals											4	4	4	4	4	4	4	4	4	4	4	4	4	4

+ Indicates animal exhibiting sign

Table 5: Time of death and clinical signs at 2101 mg/kg dose

Dosage mg/kg	Signs of reaction to treatment	Animal number and sex										Number showing effect at time after dosing														
		A856 Male					A857 Female					Hour						Day								
		1	2	3	4	5	1	2	3	4	5	1/4	1/2	1	2	4	6	2		3	4	5	6	7	8-15
																		am	pm						
2101	**Early decedents**																								
	Unconsciousness	+	+			+	+	+	+	+		7	7	6	2										
	Death	+	+			+	+	+	+	+				1	5	7	7	7	7	7	7	7	7	7	7
	Time of death (hours)	2	1			4	2	2	4	2															
	No. of animals											7	7	7	7	7	7	7	7	7	7	7	7	7	7
	Animals surviving to Day 15:																								
	None																	1	1	3	3	3	3	3	3
	Piloerection			+	+						+						2	1							
	Abdominal hunching										+						1								
	Hypoactivity			+	+						+						2	2	2						
	Unconsciousness			+	+						+	3	3	3	3	3	2								
	Staining about the muzzle			+															1						
	Ataxia				+						+						2								
	No. of animals											3	3	3	3	3	3	3	3	3	3	3	3	3	3

+ Indicates animal exhibiting sign

Table 6: Time of death and clinical signs at 2500 mg/kg dose

Dosage mg/kg	Signs of reaction to treatment	Animal number and sex										Number showing effect at time after dosing														
		A858 Male					A859 Female					Hour						Day								
		1	2	3	4	5	1	2	3	4	5	1/4	1/2	1	2	4	6	2		3	4	5	6	7	8-15
																		am	pm						
2500	**Early decedents**																								
	Unconsciousness	+	+	+	+		+	+	+	+		8	8	2											
	Hypopnea			+								1	1												
	Death	+	+	+	+		+	+	+	+				6	8	8	8	8	8	8	8	8	8	8	8
	Time of death (hours)	1	1	1	1		1	2	2	1															
	No. of animals											8	8	8	8	8	8	8	8	8	8	8	8	8	8
	Animals surviving to Day 15:																								
	None																	2	2	2	2	2	2	2	2
	Unconsciousness					+					+	2	2	2	2	2	2								
	Dyspnoea					+					+						2								
	Hypopnea					+						1	1	1	1	1									
	No. of animals											2	2	2	2	2	2	2	2	2	2	2	2	2	2

+ Indicates animal exhibiting sign

- **Necropsy findings:** See Table 7. Necropsy of early decedents invariably revealed fluid or mucoid dilatation of the gastro-intestinal tract, which was attributed to irritation. Necropsy of animals that survived the observation period revealed no consistent treatment-related effects.

Table 7: Findings at necrospy

Observation	Incidence of animals with specified pathological changes																			
	Early decendents										Animals surviving to Day 15									
	Dosage (mg/kg) administered to males					Dosage (mg/kg) administered to females					Dosage (mg/kg) administered to males					Dosage (mg/kg) administered to females				
	1247	1484	1765	2101	2500	1247	1484	1765	2101	2500	1247	1484	1765	2101	2500	1247	1484	1765	2101	2500
External																				
NSL			2	2	3		1		2	4	5	5	3	2	1	5	4		1	1
Muzzle staining					1			1	2											
Suprascapular hair loss				1																
Internal																				
NSL											5	3	3	2	1	2	4	1		1
Lung – areas of congestion on all lobes								1				2							1	
Stomach contents – fluid			2	1	3			3	3	3										
granular				1				1												
soft brown material							1													
Mucosal congestion in pylorus					1															
Duodenum contents – fluid										4										
mucoid			2	3	4		1	4	4	3										
Jejunum contents - fluid					2					2										
mucoid			2	3			1	4	4	1						2				
tympanic								1												
Ileum contents – mucoid			1	1	2			1	1	2										
Caecum contents – mucoid					1															
Cannibalised end of tail																1				
Number of animals examined	0	0	2	3	4	0	1	4	4	4	5	5	3	2	1	5	4	1	1	1

NSL No significant lesion

CONCLUSIONS

Remarks: From the observed mortality data, the acute median lethal dosage (LD_{50}) and 95% confidence interval of Dinol, calculated by probit analysis were found to be 1880 (1691 – 2120) mg/kg

DATA QUALITY

- **Reliabilities:** 1, Reliable without restriction

Remarks: Well-conducted study, similar to OECD guidelines

REFERENCES

Life Science Research, Dynol: Acute oral toxicity in the rat, Report No. 79/DSB002/565, December 1979

OTHER

- **Last Changed:** March 18, 2004
- **Order number for sorting:** 2

Remarks:

HUMAN HEALTH ENDPOINTS (NON SIDS)
13.3 SKIN IRRITATION

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Dead Sea Bromine Company Ltd

METHOD

- **Method:** U.S. Federal Register, §191.11, 1964
- **Species/strain:** Rabbit, New Zealand White
- **Concentration:** 0.5 g
- **No of animals:** 6
- **Vehicle:** Slightly moistened with distilled water
- **GLP:** No
- **Year:** 1979

Remarks: Dynol was used as supplied, slightly moistened with distilled water for easier adherence. Intact and abraded rabbit skin was tested. Occlusive dressing was used.

RESULTS

- Mildly irritating

Remarks: Upon single dermal application a barely discernible erythematous response on the abraded skin of one animal at 72 hours accounted for a Primary Irritation Index of zero. Bandage burn was recognised on 3 animals and slight exfoliation was present in one animal on day 8.

CONCLUSIONS

Remarks: A single dermal application of test substance produced a Primary Irritation Index of zero, thus placing the material into the lowest available class of 'mildly irritating' to the skin.

REFERENCES (Free Text)

Life Science Research, Dynol: Irritance to rabbit skin (US Federal Register, 1964), Report No. 79/DSB002/440, September 1979

OTHER

- Last Changed: October 3 2002
- Order number for sorting: 1

Remarks: None

HUMAN HEALTH ENDPOINTS (NON SIDS)
13.4 EYE IRRITATION

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Dead Sea Bromine Company Ltd

METHOD

- **Method:** U.S. Federal Register, §191.12, 1964
- **Species/strain:** Rabbit, New Zealand White
- **Concentration:** 100 mg
- **No of animals:** 6
- **Vehicle:** none
- **GLP:** No
- **Year:** 1979

Remarks: Dynol was used as supplied. 100 mg portions of the test material was placed in the right eye of each test animal. Ocular reaction to treatment was assessed at 24, 48 and 72 hours and seven days after installation. Scoring was carried out using the Draize system.

RESULTS

- Irritant

Remarks: Diffuse corneal opacity affecting less than a quarter of the corneal surface was exhibited by four of the test animals at the 24, 48 and 72-hour post-installation reading. One animal displayed diffuse crimson redness of the conjunctivae in addition to the corneal lesions. All animals exhibited slight conjunctivitis at the 24, 48 and 72-hour readings. This persisted to day 8 in 2 animals. Immediate pain responses upon instillation of the test material ranged from practically no initial pain to moderate initial pain.

CONCLUSIONS

Remarks: A single installation of 100 mg of test substance into the conjunctival sac of six New Zealand White rabbits provoked corneal and conjunctival lesions in four animals regarded as positive reactions under this method. Therefore the test substance was considered to elicit a positive result as an eye irritant. The positive reactions had resolved by Day 8 and may therefore be considered as reversible.

REFERENCES (Free Text)

Life Science Research, Dynol: Irritance to the rabbit eye (US Federal Register, 1964), Report No. 79/DSB/003/439, September 1979

OTHER

- Last Changed: October 3 2002
- Order number for sorting: 1

Remarks: None

HUMAN HEALTH ENDPOINTS
14.1 GENETIC TOXICITY IN VIVO

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Dow Chemical Company, Lot No.: 840429-162, Purity: approximately 78.6%, Identity of impurities: 2,2-bis (hydroxymethyl)-1-bromo-3-hydroxypropane (6.6%), 2,2-bis (bromomethyl)-1-bromo-3-hydroxypropane (6.9%), pentaerythritol (0.2%), dimers and structural isomers (7.7%).

METHOD

- **Method/guideline:** Environmental Health Research & Testing, Inc. method. 3-dose gavage protocol with the test substance administered at 24-hour intervals followed by bone marrow sampling 24 hours after third dosing.
- **Type:** Mouse Micronucleus test
- **GLP:** Yes
- **Year:** 1996
- **Species:** Mice
- **Strain:** B6C3F_1
- **Sex:** Male
- **Route of administration:** Oral, gavage
- **Doses/concentration levels:** 100, 200, 300, 400 mg/kg bw
- **Exposure period:** 4 days
- **Statistical methods:** Frequency of micronucleated cells among PCEs analyzed using a one-tailed trend test across dose groups and a t-test for pairwise comparisons of each dose group to the concurrent control.

Remarks:

- **Age at study initiation:** No data
- **No. of animals per dose:** 5
- **Vehicle:** Corn oil
- **Duration of test:** 4 days
- **Frequency of treatment:** Once every 24 hours for 3 doses
- **Sampling times and number of samples:** Sampled 24 hours after final dose
- **Control groups:** Solvent control: Corn oil;
  Positive control: Injection of 12.5 mg dimethylbenzanthracene per kg bw.
- **Clinical observations performed:** none
- **Organs examined at necropsy:** none
- **Criteria for evaluating results:** 2000 polychromatic erythrocytes scored for frequency of micronucleated cells (5 animals per dose group).
- **Criteria for selection of M.T.D.:** Dose levels were selected after a 13-week repeat dose toxicity study.

RESULTS

- **Effect on mitotic index or PCE/NCE ratio by dose level by sex:** No data
- **Genotoxic effects:** None in first trial, but there was a clear dose-related increase in micronucleated PCEs in the second trial.

- **NOAEL (NOEL)(C)/LOAEL(LOEL)(C):** No data
- **Statistical results, as appropriate:**

Remarks:

- **Mortality at each dose level by sex:** 3 mice died in the first experiment at 400 mg/kg
- **Frequency of Micronuclei:**

Dose (mg/kg)	Micronucleated Cells/1000 PCEs [a]	
	Trial 1 - Negative	Trial 2 – Positive
Positive control (12.5 mg/kg)	4.6±1.1	7.8±1.3
Solvent control	1.4±0.6	1.5±0.5
100	0.7±0.4	2.3±0.3
200	2.5±0.5	2.6±0.7
300	2.0±0.7	
400 [b]	1.2±1.2	4.8±1.2*
	P=0.220 [c]	P=0.000

* Significantly different (P<0.008) from control

[a] data presented as mean ± standard error; PCE = polychromatic erythrocyte

[b] Only 2 mice survived this group in Trial 1

[c] Trend Test

- **Description, severity, time of onset and duration of clinical signs at each dose level and sex:** no data
- **Bodyweight changes by dose and sex:** No data
- **Food/water consumption by dose and sex:** No data

CONCLUSIONS

Remarks: Results of the first trial were negative. In the second trial, however, there was a clear dose-related increase in micronucleated PCEs. Because the positive response was not reproduced the results were concluded to be equivocal.

DATA QUALITY

- Reliabilities: 2, Reliable with restrictions

Remarks:

REFERENCES (Free Text)

NTP Technical Report on the Toxicology and Carcinogenesis Studies of 2,2-bis(bromomethyl)-1,3-propanediol (FR-1138, CAS No. 3296-90-0) in F344/N Rats and B6C3F1 Mice (Feed Studies), NTP TR 452, National Toxicology Program, May 1996

OTHER

- **Last Changed:** October 3 2002
- **Order number for sorting:** 1

Remarks:

HUMAN HEALTH ENDPOINTS
14.2 GENETIC TOXICITY IN VIVO

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Dow Chemical Company, Lot No.: 840429-162, Purity: approximately 78.6%, Identity of impurities: 2,2-bis (hydroxymethyl)-1-bromo-3-hydroxypropane (6.6%), 2,2-bis (bromomethyl)-1-bromo-3-hydroxypropane (6.9%), pentaerythritol (0.2%), dimers and structural isomers (7.7%).

METHOD

- **Method/guideline:** Environmental Health Research & Testing, Inc. method. Single intraperitoneal injection followed by bone marrow sampling 48 hours after dosing.
- **Type:** Mouse Micronucleus test
- **GLP:** Yes
- **Year:** 1996
- **Species:** Mice
- **Strain:** B6C3F_1
- **Sex:** Male/Female
- **Route of administration:** Intraperitoneal injection
- **Doses/concentration levels:** 150, 300, 600 mg/kg bw
- **Exposure period:** 2 days
- **Statistical methods:** Data analyzed by the Cochran-Armitage trend test and pairwise comparison of dose groups to the corresponding negative controls were made using a t-test.

Remarks:

- **Age at study initiation:** No data
- **No. of animals per dose:** 3 or 4
- **Vehicle:** Corn oil
- **Duration of test:** 2 days
- **Frequency of treatment:** Single dose
- **Sampling times and number of samples:** Sampled 48 hours after dosing
- **Control groups:** Solvent control: Corn oil;
  Positive control: Injection of 200 mg/kg urethane.
- **Clinical observations performed:** none
- **Organs examined at necropsy:** none
- **Criteria for evaluating results:** 1000 polychromatic erythrocytes scored for frequency of micronucleated cells.
- **Criteria for selection of M.T.D.:** Dose levels were selected after a 13-week repeat dose toxicity study and oral bone marrow micronucleus study.

RESULTS

- **Effect on mitotic index or PCE/NCE ratio by dose level by sex:** No data
- **Genotoxic effects:** Male mice in all three dose groups showed a 2-fold increase in the frequency of micronucleated PCEs. Trend test was not significant due to the similarity in responses and pairwise analyses were also not significant. Female response was stronger

(2.5-fold increase over controls at highest dose) and was directly related to increasing doses of test substance

- **NOAEL (NOEL)(C)/LOAEL(LOEL)(C):** No data
- **Statistical results, as appropriate:**

Remarks:

- **Mortality at each dose level by sex:** No deaths reported
- **Frequency of Micronuclei:**

Dose (mg/kg)	Micronucleated Cells/1000 PCEs [a]	
	Male	Female
Positive control (200 mg/kg)	16.4±2.2 (3) [b]	12.1±0.9 (4)
Solvent control	1.5±0.3 (4)	2.0±0.4 (4)
150	3.2±0.8* (4)	2.7±1.1 (4)
300	3.0±0.7* (4)	3.6±0.9* (3)
600	3.0±1.0* (3)	5.2±0.5* (4)
	P=0.150 [c]	P=0.003

* Significantly different (P<0.008) from control

[a] Data presented as mean ± standard error; PCE = polychromatic erythrocyte

[b] Number of mice in parentheses

[c] Trend Test

- **Description, severity, time of onset and duration of clinical signs at each dose level and sex:** no data
- **Bodyweight changes by dose and sex:** No data
- **Food/water consumption by dose and sex:** No data

CONCLUSIONS

Remarks: The results of this experiment provide evidence of the ability of the test substance to induce micronuclei in bone marrow cells of female mice. The results were consistent with the stronger response observed in female mice in the 13-week feed study.

DATA QUALITY

- Reliabilities: 2, Reliable with restrictions

Remarks:

REFERENCES (Free Text)

NTP Technical Report on the Toxicology and Carcinogenesis Studies of 2,2-bis(bromomethyl)-1,3-propanediol (FR-1138, CAS No. 3296-90-0) in F344/N Rats and B6C3F1 Mice (Feed Studies), NTP TR 452, National Toxicology Program, May 1996

OTHER

- **Last Changed:** October 3 2002
- **Order number for sorting:** 2

Remarks:

HUMAN HEALTH ENDPOINTS
14.3 GENETIC TOXICITY IN VIVO

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Dow Chemical Company, Lot No.: 840429-162, Purity: approximately 78.6%, Identity of impurities: 2,2-bis (hydroxymethyl)-1-bromo-3-hydroxypropane (6.6%), 2,2-bis (bromomethyl)-1-bromo-3-hydroxypropane (6.9%), pentaerythritol (0.2%), dimers and structural isomers (7.7%).

METHOD

- **Method/guideline:** MacGregor, et al (1990). Blood samples were obtained from animals at the end of the 13 week repeat oral dose toxicity study.
- **Type:** Mouse Peripheral Blood Micronucleus test
- **GLP:** Yes
- **Year:** 1996
- **Species:** Mice
- **Strain:** B6C3F_1
- **Sex:** Male/Female
- **Route of administration:** Oral
- **Doses/concentration levels:** 0, 625, 1250, 2500, 5000, 10000 ppm
- **Exposure period:** 13 weeks
- **Statistical methods:** Log transformation of NCE data, testing for normality by the Shapiro-Wilk test, and testing for heterogeneity of variance by Cochran's test were performed before statistical analyses. The frequency of micronucleated cells among NCEs was analyzed by analysis of variance using the SAS GLM procedure. The NCE data for each dose group were compared with the concurrent solvent control using a Student's t-test.

Remarks:

- **Age at study initiation:** 6 weeks
- **No. of animals per dose:** 7-10
- **Vehicle:** Feed
- **Duration of test:** 13 weeks
- **Frequency of treatment:** Continuous, ad libitum
- **Sampling times and number of samples:** 1 sample taken at end of study
- **Control groups:** Feed control
- **Clinical observations performed:** none
- **Organs examined at necropsy:** none
- **Criteria for evaluating results:** 10000 normochromatic erythrocytes scored for frequency of micronucleated cells.
- **Criteria for selection of M.T.D.:**

RESULTS

- **Actual dose received:** 100, 200, 500, 1300, 3000 mg/kg bw (male)
  140, 300, 600, 1200, 2900 mg/kg bw (female)
- **Effect on mitotic index or PCE/NCE ratio by dose level by sex:** No data

- **Genotoxic effects:** Significant increases in micronucleated normochromatic erythrocytes were observed in the 2 highest dose groups of male mice (5000 and 10000 ppm) and the 3 highest dose groups of female mice (2500 to 10000 ppm).
- **NOEL:** 625 ppm (males, not achieved for females)
- **LOEL:** 1250 ppm (males), 625 ppm (females)
- **Statistical results, as appropriate:**

Remarks:

- **Mortality at each dose level by sex:** one female (control), 2 males & 1 female (625 ppm), 1 female (1250 ppm), 1 female (2500 ppm), 1 female (5000 ppm), 3 males (10000 ppm) died during the study.

- **Frequency of Micronuclei:**

Dose (ppm)	Micronucleated NCEs/1000 Cells [a]	
	Male	Female
0	2.36±0.17 (10) [b]	1.46±0.26 (9)
625	2.28±0.29 (8)	1.86±0.30 (9)
1250	2.55±0.18 (10)	1.86±0.22 (9)
2500	2.98±0.21 (10)	2.72±0.32 [c] (9)
5000	3.80±0.19[c] (10)	4.26±0.47 [c] (9)
10000	9.30±1.26 [c]	11.81±0.54 [c] (9)
	P<0.001 [d]	P<0.001

[a] Data presented as mean ± standard error; NCE = normochromatic erythrocyte

[b] Number of mice in parentheses

[c] Significant response by pairwise comparison to control

[d] Trend test

- **Description, severity, time of onset and duration of clinical signs at each dose level and sex:** Clinical findings included abnormal posture and hypoactivity in 10000-ppm male and female mice.

- **Bodyweight changes by dose and sex:** The final mean body weights and body weight gains of males and females receiving 1250, 2500, 5000 or 10000-ppm and of females receiving 625 ppm were significantly lower than those of the controls.

- **Food/water consumption by dose and sex:** Feed consumption by exposed mice was generally higher than that by controls throughout the study.

CONCLUSIONS

Remarks: The test substance was shown to be genotoxic in vivo

DATA QUALITY

- Reliabilities: 2, Reliable with restrictions

Remarks:

REFERENCES (Free Text)

NTP Technical Report on the Toxicology and Carcinogenesis Studies of 2,2-bis(bromomethyl)-1,3-propanediol (FR-1138, CAS No. 3296-90-0) in F344/N Rats and B6C3F1 Mice (Feed Studies), NTP TR 452, National Toxicology Program, May 1996

MacGregor, J.T., et al, The in vivo erythrocyte micronucleus test: Measurement at steady state increases assay efficiency and permits integration with toxicity studies. Fundam. Appl. Toxicol., 14, 513-522, 1990

OTHER

- **Last Changed:** March 26, 2003
- **Order number for sorting:** 3

Remarks:

HUMAN HEALTH ENDPOINTS
15.1 GENETIC TOXICITY IN VITRO

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Bromine Compounds Ltd.
Substance: FR-522 Purified (2,2-bis(bromomethyl)-1,3-propanediol), Purity: 99.5%, Lot No.: 616-521-11

METHOD

- **Method:** OECD 471, 92/69/EEC Method B14, US EPA, Method: HG-Gene Muta – S. typhimurium: The Salmonella typhimurium reverse mutation assay, 1984
- **Test Type:** Reverse Mutation Assay
- **System of testing:** Bacterial
- **GLP:** Yes
- **Year:** 1995
- **Species/Strain:** Salmonella typhimurium strains TA98, TA100, TA1535 and TA1537
- **Metabolic activation:** S9-mix, Rat liver cells, Aroclor induced
  S9-mix, Hamster liver cells, uninduced
- **Concentrations tested:** 50, 150, 500, 1500, 5000 µg/plate
- **Statistical Methods:** None

Remarks:

- **Test Design**
  - **Number of replicates:** 3
  - **Positive Controls:** N-ethyl-N'-nitroso-N-nitrosoguanidine (TA100, TA1535, -S9)
    9-aminoacridine (TA1537, -S9)
    2-nitrofluorene (TA98, -S9)
    2-aminoanthracene (TA98, TA100, TA 1535, TA1537, +S9)
    Congo red (TA98, +S9)
  - **Negative Control:** Solvent vehicle
- **Solvent:** Dimethylsulfoxide

RESULTS

- **Result:** Positive
- **Cytotoxic concentration**
  - **With metabolic activation:** > 5000 µg/plate
  - **Without metabolic activation:** > 5000 µg/plate
- **Genotoxic effects**
  - **With metabolic activation:** Large dose-related increases in revertant colony numbers were observed with strain TA1535. These increases were observed only in the presence of hamster S-9

mix and were largest in the presence of the 30% mix. A smaller increase was also observed with strain TA100 in the presence of 30% hamster S-9 mix.

- **Without metabolic activation:** None

- **Statistical results:**

Strain	Dose level (µg/plate)	Revertants/Plate [a]							
		Test 1				Test 2			
		-S9	+10% Rat S9	+10% Hamster S9	+30% Hamster S9	-S9	+10% Rat S9	+10% Hamster S9	+30% Hamster S9
TA1535	5000	17±4.0	17±1.7	67±14.6	243±28.8	16±3.6	15±3.0	85±20.0	217±15
	1500	15±5.5	11±1.0	27±0.6	90±4.5	15±2.1	15±4.0	35±5.3	92±5.7
	500	12±4.5	14±5.8	17±4.6	52±21.9	17±4.6	13±4.6	21±5.5	71±0.6
	150	16±2.5	12±2.3	17±5.0	24±5.5	13±4.0	12±3.6	20±6.1	37±6.4
	50	16±6.1	15±3.6	15±3.6	22±5.1	17±1.2	12±0.0	18±2.3	18±5.6
	0	13±1.2	14±4.5	15±4.5	15±3.2	15±1.5	13±3.1	15±4.9	13±3.8
	Solvent	18±5.5	12±1.7	16±5.2	15±3.6	15±4.5	19±3.2	16±3.1	17±0.7
	ENNG (5.0)	1361±20.1				1485±16.8			
	AA (2.0)		88±14.2	157±4.7	137±18.6		185±5.3	202±18.0	203±16.8
TA1537	5000	7±0.6	6±2.3	8±3.5	13±2.1	14±3.6	11±2.0	10±4.2	12±3.1
	1500	9±3.2	8±3.6	11±1.0	11±3.5	14±2.0	10±4.2	16±1.5	11±3.2
	500	6±0.0	7±4.7	10±2.6	10±5.1	16±2.6	11±1.0	11±2.6	11±2.9
	150	5±1.7	9±2.5	10±3.1	10±2.9	11±2.5	12±4.0	13±2.1	11±4.0
	50	8±3.2	11±1.2	10±3.5	10±2.5	15±4.7	12±3.6	15±2.0	11±2.6
	0	8±1.7	14±2.6	9±3.1	13±1.5	8±3.0	9±1.0	13±4.0	11±2.6
	Solvent	10±0.6	12±3.8	12±1.7	11±2.1	11±5.0	14±1.7	11±1.2	12±1.5
	9 AC (80.8)	X				X			
	AA (2.0)		67±15.5	459±22.5	383±25.1		163±15.3	235±51.9	257±38.2
TA98	5000	21±2.0	26±4.7	31±2.6	28±5.1	27±0.0	28±6.4	24±1.7	29±4.0
	1500	21±1.5	26±3.8	23±3.6	26±5.5	28±5.5	26±4.9	26±4.0	29±0.6
	500	20±3.6	26±3.5	25±3.8	26±3.5	25±5.0	24±3.8	28±4.2	22±3.5
	150	22±3.8	31±2.9	26±2.5	25±1.7	21±1.2	27±1.5	25±1.7	26±4.0
	50	18±4.0	25±6.0	24±4.0	27±2.6	27±2.5	28±1.0	22±2.6	24±4.2
	0	24±1.5	28±1.5	23±2.3	26±4.6	22±0.6	25±4.7	26±4.5	28±3.1
	Solvent	23±0.6	29±2.1	23±3.2	25±2.1	24±2.1	25±6.4	25±4.4	27±0.0
	NF (1.0)	338±7.0				275±6.2			
	AA (0.5)		152±16.8	122±64.4	167±26.8		232±19.3	237±17.3	194±9.2
	CR (200.0)		20±2.5	34±11.5	105±13.6		43±9.6	64±6.2	218±17.4
TA100	5000	116±4.2	125±8.2	136±19.1	271±28.0	122±5.7	122±11.8	138±26.2	242±46.4
	1500	130±4.2	130±7.8	148±17.8	175±7.0	120±3.6	119±11.0	136±6.7	201±6.6
	500	116±8.6	134±7.5	143±2.6	140±11.3	132±11.6	119±5.8	138±10.7	176±12.7
	150	130±5.0	122±2.0	127±2.6	141±7.0	124±10.4	133±12.3	128±13.1	175±20.0
	50	130±3.6	119±14.9	124±11.9	152±11.1	109±5.1	138±18.7	128±17.0	128±6.1
	0	125±8.1	140±13.1	144±13.8	150±5.6	122±17.1	127±14.5	137±15.9	136±11.5
	Solvent	124±2.3	118±13.0	119±8.7	143±11.5	115±11.1	123±15.9	143±3.5	137±18.9
	ENNG (3.0)	654±50.3				1568±52.8			
	AA (1.0)		460±3.6	886±21.6	485±123.8		444±24.1	723±41.6	650±121.3

[a] Data presented as mean ± standard deviation

ENNG N-ethyl-N'-nitro-N-nitrosoguanidine, 9 AC 9-aminoacridine, NF 2-nitrofluorene, AA 2-aminoanthracene, CR Congo red

X Too many colonies to count accurately

Remarks: None

CONCLUSIONS

Remarks: It is concluded that, when tested in dimethylsulfoxide, the test substance shows no evidence of mutagenic activity in the absence or presence of rat S-9 mix. The test substance shows clear evidence of mutagenic activity with strains TA1535 and TA100 in the presence of hamster S-9 mix.

DATA QUALITY

- **Reliabilities:** 1, Reliable without restrictions

Remarks: Study conducted under GLP to OECD/EC/EPA test guidelines by Huntingdon Life Sciences.

REFERENCES

Huntingdon Life Sciences Ltd., FR-522 Purified, Bacterial Mutation Assay, February 1996

OTHER

- **Last Changed:** October 3 2002
- **Order number for sorting:** 1

Remarks: S-9 is usually only prepared from a rat previously treated with a compound (Aroclor) known to induce a high level of enzymic activity. However in this study S-9 mix using liver fraction from uninduced Syrian hamsters was also used as it was suspected that the test substance may be metabolised to a mutagen by hamster S-9 but not by rat.

HUMAN HEALTH ENDPOINTS
15.2 GENETIC TOXICITY IN VITRO

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Bromine Compounds Ltd.
Substance: FR-522 (2,2-bis(bromomethyl)-1,3-propanediol), Purity: 98.63%, Lot No.: 953038

METHOD

- **Method:** OECD 471, 92/69/EEC Method B14, US EPA, Method: HG-Gene Muta– S. typhimurium: The Salmonella typhimurium reverse mutation assay, 1984
- **Test Type:** Reverse Mutation Assay
- **System of testing:** Bacterial
- **GLP:** Yes
- **Year:** 1996
- **Species/Strain:** Salmonella typhimurium strains TA98, TA100, TA1535 and TA1537
- **Metabolic activation:** S9-mix, Rat liver cells, Aroclor induced
  S9-mix, Hamster liver cells, uninduced
- **Concentrations tested:** 50, 150, 500, 1500, 5000 µg/plate
- **Statistical Methods:** None

Remarks:

- **Test Design**

  - **Number of replicates:** 3
  - **Positive Controls:** N-ethyl-N'-nitroso-N-nitrosoguanidine (TA100, TA1535, -S9)
    9-aminoacridine (TA1537, -S9)
    2-nitrofluorene (TA98, -S9)
    2-aminoanthracene (TA98, TA100, TA 1535, TA1537, +S9)
    Congo red (TA98, +S9)
  - **Negative Control:** Solvent vehicle

- **Solvent:** Dimethylsulfoxide

RESULTS

- **Result:** Positive
- **Cytotoxic concentration**

  - **With metabolic activation:** > 5000 µg/plate
  - **Without metabolic activation:** > 5000 µg/plate

- **Genotoxic effects**
  - **With metabolic activation:** Fairly large dose-related increases in revertant colony numbers were observed with strains TA1535 and TA 100. These increases were observed only in the presence of hamster S-9 mix and were largest in the presence of the 30% mix.

- **Without metabolic activation:** None

- **Statistical results:**

Strain	Dose level (µg/plate)	Revertants/Plate [a]							
		Test 1				Test 2			
		-S9	+10% Rat S9	+10% Hamster S9	+30% Hamster S9	-S9	+10% Rat S9	+10% Hamster S9	+30% Hamster S9
TA1535	5000	13±4.9	15±4.6	46±11.8	108±15.9	10±4.6	11±4.2	33±6.0	177±16.2
	1500	17±3.1	18±3.6	53±8.2	73±5.9	11±4.7	12±4.0	30±3.6	49±17.1
	500	12±2.3	17±4.6	16±3.1	29±13.1	13±2.3	8±3.5	19±11.0	37±1.0
	150	11±3.0	12±1.5	18±2.6	29±5.2	14±5.5	14±4.5	17±4.5	10±4.0
	50	12±5.3	13±3.0	11±2.1	12±6.7	12±3.6	16±2.1	16±4.6	12±2.5
	0	11±0.6	15±2.0	14±4.6	14±5.3	8±1.2	9±2.5	13±5.0	11±5.3
	Solvent	13±3.5	16±4.6	11±3.5	14±3.8	10±1.5	16±3.8	11±3.5	11±1.0
	ENNG (5.0)	1192±83.4				1455±49.7			
	AA (2.0)		77±11.0	116±68.5	160±5.6		87±13.7	91±15.0	152±11.0
TA1537	5000	12±5.6	10±6.7	14±2.1	9±3.1	9±0.6	8±1.2	8±1.2	10±1.0
	1500	14±4.2	12±4.6	11±0.6	13±5.3	7±1.0	7±1.5	7±1.5	9±3.5
	500	11±3.2	12±1.0	12±2.6	15±2.6	9±4.4	9±2.1	7±6.1	10±2.0
	150	15±4.6	12±2.1	16±3.5	13±3.1	12±4.7	9±4.0	11±1.0	12±1.5
	50	12±6.4	14±3.6	15±3.5	11±4.9	15±4.4	14±5.0	10±2.1	15±1.7
	0	11±1.0	11±0.0	14±1.2	11±2.9	9±1.0	8±1.2	10±4.0	8±4.9
	Solvent	9±1.0	13±2.9	12±1.2	11±3.1	8±0.6	8±0.6	10±3.1	9±2.6
	9 AC (80.0)	X				X			
	AA (2.0)		42±5.8	195±12.1	160±41.9		71±4.7	142±10.8	163±25.1
TA98	5000	22±6.6	26±5.9	26±5.5	26±4.5	25±1.7	27±1.2	22±2.6	26±4.0
	1500	27±1.7	30±5.7	30±6.0	31±4.9	23±2.6	22±3.5	24±3.6	29±3.2
	500	22±5.6	29±5.1	23±2.1	23±3.5	22±4.7	27±5.5	23±4.6	28±4.5
	150	28±3.2	30±3.2	23±7.0	20±4.2	22±2.5	25±3.2	22±0.6	25±1.7
	50	26±2.5	29±3.5	23±4.0	33±2.6	23±5.3	23±5.9	26±2.3	27±0.6
	0	22±4.0	27±4.2	22±4.4	29±4.7	18±2.0	22±1.0	24±2.1	28±3.6
	Solvent	28±3.1	27±4.5	26±2.0	28±3.2	18±1.7	25±4.0	20±1.0	28±4.2
	NF (1.0)	146±23.4				274±8.1			
	AA (0.5)		101±16.7	419±9.3	392±82.1		170±30.8	376±30.5	397±14.5
	CR (200.0)		33±3.2	39±3.1	127±13.3		32±2.3	57±1.7	181±11.7
TA100	5000	125±11.8	126± -	159±14.1	178±24.3	100±8.1	121±5.0	146±12.8	239±50.0
	1500	129±13.7	135±2.6	137±4.9	182±14.4	98±22.4	113±6.6	153±3.5	186±1.5
	500	93±3.8	115±10.8	125±18.5	131±12.7	110±9.5	106±15.6	115±6.6	134±9.9
	150	98±7.2	115±10.1	129±8.1	148±13.7	118±14.6	133±10.3	140±14.5	142±11.2
	50	115±13.4	131±15.7	119±5.0	133±3.6	105±7.1	112±6.1	108±12.9	128±8.1
	0	110±16.5	112±14.0	129±19.3	124±12.2	116±18.6	115±12.5	103±14.7	128±10.7
	Solvent	105±3.0	110±6.4	126±9.3	129±7.1	101±9.0	124±1.0	127±13.9	123±8.0
	ENNG (3.0)	825±115.8				1426±34.8			
	AA (1.0)		333±109.3	837±146.7	613±313.8		324±33.0	979±127.3	519±122.4

[a] Data presented as mean ± standard deviation

ENNG N-ethyl-N'-nitro-N-nitrosoguanidine, 9 AC 9-aminoacridine, NF 2-nitrofluorene, AA 2-aminoanthracene, CR Congo red

X Too many colonies to count accurately

Remarks: None

CONCLUSIONS

Remarks: It is concluded that, when tested in dimethylsulfoxide, the test substance shows no evidence of mutagenic activity in the absence or presence of rat S-9 mix. The test substance shows clear evidence of mutagenic activity with strains TA1535 and TA100 in the presence of hamster S-9 mix.

DATA QUALITY

- **Reliabilities:** 1, Reliable without restrictions

Remarks: Study conducted under GLP to OECD/EC/EPA test guideline by Huntingdon Life Sciences.

REFERENCES

Huntingdon Life Sciences Ltd., FR-522, Bacterial Mutation Assay, January 1996

OTHER

- **Last Changed:** October 3 2002
- **Order number for sorting:** 2

Remarks: S-9 is usually only prepared from a rat previously treated with a compound (Aroclor) known to induce a high level of enzymic activity. However in this study S-9 mix using liver fraction from uninduced Syrian hamsters was also used as it was suspected that the test substance may be metabolised to a mutagen by hamster S-9 but not by rat.

HUMAN HEALTH ENDPOINTS
15.3 GENETIC TOXICITY IN VITRO

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Bromine Compounds Ltd.
Substance: FR-522C (2,2-bis(bromomethyl)-1,3-propanediol), Purity: 88.4%, Lot No.: 1138

METHOD

- **Method:** OECD 471, 92/69/EEC Method B14, U S EPA, Method: HG-Gene Muta– S. typhimurium: The Salmonella typhimurium reverse mutation assay, 1984
- **Test Type:** Reverse Mutation Assay
- **System of testing:** Bacterial
- **GLP:** Yes
- **Year:** 1996
- **Species/Strain:** Salmonella typhimurium strains TA98, TA100, TA1535 and TA1537
- **Metabolic activation:** S9-mix, Rat liver cells, Aroclor induced
  S9-mix, Hamster liver cells, uninduced
- **Concentrations tested:** 50, 150, 500, 1500, 5000 µg/plate
- **Statistical Methods:** None

Remarks:

- **Test Design**

  - **Number of replicates:** 3
  - **Positive Controls:** N-ethyl-N'-nitroso-N-nitrosoguanidine (TA100, TA1535, -S9)
    9-aminoacridine (TA1537, -S9)
    2-nitrofluorene (TA98, -S9)
    2-aminoanthracene (TA98, TA100, TA 1535, TA1537, +S9)
    Congo red (TA98, +S9)
  - **Negative Control:** Solvent vehicle

- **Solvent:** Dimethylsulfoxide

RESULTS

- **Result:** Positive
- **Cytotoxic concentration**

  - **With metabolic activation:** > 5000 µg/plate
  - **Without metabolic activation:** > 5000 µg/plate

- **Genotoxic effects**
  - **With metabolic activation:** Large dose-related increases in revertant colony numbers were observed with strains TA1535 and TA 100. These increases were observed only in the presence of hamster S-9 mix.
  - **Without metabolic activation:** None

- **Statistical results:**

Strain	Dose level (µg/plate)	Revertants/Plate [a]							
		Test 1				Test 2			
		-S9	+10% Rat S9	+10% Hamster S9	+30% Hamster S9	-S9	+10% Rat S9	+10% Hamster S9	+30% Hamster S9
TA1535	5000	10±3.5	14±2.1	97±86	250±32.4	13±3.2	11±3.8	46±8.1	122±8.5
	1500	15±5.6	18±2.6	86±14.7	179±7.5	10±1.0	10±3.0	39±6.7	82±20.8
	500	12±3.0	15±2.1	57±6.6	118±15.1	13±1.2	14±4.2	38±8.5	65±28.6
	150	12±3.5	12±4.4	28±8.5	48±11.6	13±1.2	14±-	14±6.1	38±11.6
	50	8±1.7	16±4.4	30±9.1	31±1.5	17±3.2	13±3.1	17±2.1	19±1.5
	0	11±0.6	15±2.0	14±4.6	14±5.3	8±1.2	9±2.5	13±5.0	11±5.3
	Solvent	13±3.5	16±4.6	11±3.5	14±3.8	10±1.5	16±3.8	11±3.5	11±1.0
	ENNG (5.0)	1192±83.4				1455±49.7			
	AA (2.0)		77±11.0	116±68.5	160±5.6		87±13.7	91±15.0	152±11.0
TA1537	5000	9±1.7	13±4.2	9±3.1	12±2.1	11±1.7	14±7.1	14±4.4	10±2.5
	1500	10±0.6	9±3.5	12±4.0	9±2.0	11±3.6	12±4.0	10±4.4	11±1.0
	500	12±1.7	10±3.1	14±2.3	11±3.5	13±4.7	10±4.0	11±5.5	13±4.5
	150	13±2.3	8±1.2	9±2.5	13±5.5	9±1.7	12±2.9	10±4.0	13±3.5
	50	13±5.5	8±3.2	14±2.1	16±3.1	12±2.5	12±2.6	14±3.2	12±4.2
	0	11±1.0	11±0.0	14±1.2	11±2.9	9±1.0	8±1.23	10±4.0	8±4.9
	Solvent	9±1.0	13±2.9	12±1.2	11±3.1	8±0.6	8±0.6	10±3.1	9±2.6
	9 AC (80.8)	X				X			
	AA (2.0)		42±5.8	195±12.1	160±41.9		71±4.7	142±10.8	163±25.1
TA98	5000	15±5.0	24±2.1	24±5.6	31±1.0	20±2.9	20±5.7	24±7.6	26±3.5
	1500	21±5.7	31±2.5	24±1.0	25±7.8	20±3.5	23±3.2	24±2.9	25±3.6
	500	19±1.0	26±3.6	21±4.6	30±4.6	26±3.2	24±5.0	22±0.6	27±2.6
	150	18±1.2	27±5.1	27±6.8	20±3.6	20±2.1	28±1.7	26±6.5	25±7.0
	50	18±1.0	23±7.0	27±3.5	23±5.2	27±3.5	28±7.1	25±0.6	22±4.0
	0	22±4.0	27±4.2	22±4.4	29±4.7	18±2.0	22±1.0	24±2.1	28±3.6
	Solvent	28±3.1	27±4.5	26±2.0	28±3.2	18±1.7	25±4.0	20±1.0	28±4.2
	NF (1.0)	146±23.4				274±8.1			
	AA (0.5)		101±16.7	419±9.3	392±82.1		170±30.8	376±30.5	397±14.5
	CR (200.0)		33±3.2	39±3.1	127±13.3		32±2.3	57±1.7	181±11.7
TA100	5000	123±3.8	138±11.7	148±12.0	244±16.9	130±14.5	124±4.4	138±10.5	228±9.8
	1500	105±9.5	122±3.5	159±18.9	200±40.8	117±13.7	132±16.6	179±14.6	208±21.5
	500	107±15.5	134±6.5	142±9.5	157±4.0	113±10.0	123±6.7	140±3.2	161±9.8
	150	110±6.1	120±4.0	129±6.5	134±6.1	108±4.0	124±15.9	118±9.0	146±2.1
	50	115±4.0	129±3.6	141±8.5	138±25.5	117±11.0	134±5.3	130±12.3	137±4.7
	0	110±16.5	112±14.0	129±19.3	124±12.2	116±18.6	115±12.5	103±14.8	128±10.7
	Solvent	105±3.0	110±6.4	126±9.3	129±7.1	101±9.0	124±1.0	127±13.9	123±8.0
	ENNG (3.0)	825±115.8				1426±34.8			
	AA (1.0)		333±109.3	837±146.7	613±313.8		324±33.0	979±127.3	519±122.4

[a] Data presented as mean ± standard deviation

ENNG N-ethyl-N'-nitro-N-nitrosoguanidine, 9 AC 9-aminoacridine, NF 2-nitrofluorene, AA 2-aminoanthracene, CR Congo red

X Too many colonies to count accurately

Remarks: None

CONCLUSIONS

Remarks: It is concluded that, when tested in dimethylsulfoxide, the test substance shows no evidence of mutagenic activity in the absence or presence of rat S-9 mix. The test substance shows clear evidence of mutagenic activity with strains TA1535 and TA100 in the presence of hamster S-9 mix.

DATA QUALITY

- **Reliabilities:** 1, Reliable without restrictions

Remarks: Study conducted under GLP to OECD/EC/EPA test guidelines by Huntingdon Life Sciences.

REFERENCES

Huntingdon Life Sciences Ltd., FR-522C, Bacterial Mutation Assay, January 1996

OTHER

- **Last Changed:** October 3 2002
- **Order number for sorting:** 3

Remarks: S-9 is usually only prepared from a rat previously treated with a compound (Aroclor) known to induce a high level of enzymic activity. However in this study S-9 mix using liver fraction from uninduced Syrian hamsters was also used as it was suspected that the test substance may be metabolised to a mutagen by hamster S -9 but not by rat.

HUMAN HEALTH ENDPOINTS
15.4 GENETIC TOXICITY IN VITRO

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Radian Corporation
Purity: Technical grade, approximately 79% pure

METHOD

- **Method:** similar to OECD 471.
- **Test Type:** Reverse Mutation Assay
- **System of testing:** Bacterial
- **GLP:** No data
- **Year:** 1986 (Expt 1, Mortelmans, et al), 1992 (Expt 2, Zeiger, et al)
- **Species/Strain:** Salmonella typhimurium strains TA98, TA100, TA1535 and TA1537
- **Metabolic activation:** S9-mix, Rat liver cells, Aroclor induced
  S9-mix, Hamster liver cells, Aroclor induced
- **Concentrations tested:** Expt 1: 0, 33, 100, 333, 1000, 3333, 10000 µg/plate
  Expt 2: 0, 10, 33, 100, 333, 1000, 1666, 3333, 6666 µg/plate
- **Statistical Methods:** None

Remarks:

- **Test Design**

  - **Number of replicates:** 3
  - **Positive Controls:** Sodium azide (TA100, TA1535, -S9)
    9-aminoacridine (TA1537, -S9)
    4-nitro-o-phenylenediamine (TA98, -S9)
    2-aminoanthracene (TA98, TA100, TA 1535, TA1537, +S9)
  - **Negative Control:** Solvent vehicle

- **Solvent:** No data

RESULTS

- **Result:** Positive
- **Cytotoxic concentration**

  - **With metabolic activation:** > 10000 µg/plate (Expt 1), > 6666 µg/plate (Expt 2)
  - **Without metabolic activation:** > 10000 µg/plate (Expt 1), > 6666 µg/plate (Expt 2)

- **Genotoxic effects**

  - **With metabolic activation:**
    Experiment 1: Weakly positive with TA100 in the presence of 10% hamster S9 (not reproduced in second test). Negative with 10% rat S9 mix with all strains.

Experiment 2: Positive with TA100 in the presence of 30% hamster S9 mix. Negative with 30% rat S9 mix with all strains.

- **Without metabolic activation:**
  Experiment 1: None
  Experiment 2: none

- **Statistical results:**

Expt 1 (Mortelmans, et al)

Strain	Dose (µg/plate)	Revertants/Plate [a]					
		-S9		+10% hamster S9		+10% rat S9	
		Trial 1	Trial 2	Trial 1	Trial 2	Trial 1	Trial 2
TA100	0	69±3.8	83±3.5	77±4.2	94±9.0	81±3.2	76±4.6
	10		64±6.0				
	33	76±1.9	59±6.0				
	100	64±5.3	56±3.5	88±6.2	105±11.3	93±7.3	85±4.8
	333	55±5.0	61±4.0	105±7.5	93±4.0	85±7.3	76±4.8
	1000	toxic	56±2.5	112±7.2	107±9.5	93±1.2	79±6.0
	3333	toxic		126±0.0	106±8.5	82±6.9	85±7.0
	10000			toxic	110±10.0	toxic	71±8.4
Trial Summary		Negative	Negative	Weakly Positive	Negative	Negative	Negative
Positive Control		625±45.2	429±44.2	1468±2.9	1171±157.7	1362±92.4	1044±44.5
TA1535	0	6±0.9	7±0.7	6±0.6	10±0.9	8±0.3	6±2.0
	10						
	33	4±0.9	6±0.6				
	100	5±1.9	9±1.5	10±1.7	10±1.3	10±2.1	10±2.0
	333	2±1.0	6±0.6	17±2.4	14±1.9	11±2.2	10±1.2
	1000	3±1.2	10±1.8	14±3.2	10±4.5	10±0.3	10±0.3
	3333	2±0.7	toxic	16±3.0	16±1.8	12±1.7	10±0.9
	10000			12±1.2	14±1.9	10±1.8	11±0.7
Trial Summary		Negative	Negative	Equivocal	Negative	Negative	Negative
Positive Control		471±110.0	488±98.5	128±3.9	113±15.5	280±31.9	71±11.6
TA1537	0	2±0.9	6±0.6	6±1.9	10±1.5	7±0.9	10±1.5
	33	1± 0.3	4±1.2				
	100	1±0.6	8±0.9	5±0.9	7±1.2	5±1.5	6±1.2
	333	0±0.3	7±1.8	3±0.3	7±1.0	6±2.1	10±1.5
	1000	0±0.0	3±2.0	6±1.5	10±2.1	2±0.9	9±0.3
	3333	1±0.3	3±3.0	2±1.0	9±1.2	2±0.9	7±0.6
	10000			2±0.9	toxic	2±0.6	7±1.5
Trial Summary		Negative	Negative	Negative	Negative	Negative	Negative
Positive Control		432±12.9	55±6.8	74±3.5	52±7.0	58±3.0	71±15.8
TA98	0	21±2.2	18±4.0	11±0.9	21±3.2	10±0.6	23±1.7
	10		11±1.5				
	33	9±1.5	12±1.2				
	100	12±2.4	11±1.2	15±1.8	14±4.4	12±1.2	20±0.3
	333	7±1.3	12±1.5	13±1.8	19±4.6	12±1.0	20±1.2
	1000	toxic	8±0.3	10±2.1	22±1.2	13±3.4	19±2.3
	3333	toxic		14±0.6	23±1.9	13±1.2	20±3.8
	10000			3±0.9	19±3.5	8±1.2	21±1.2
Trial Summary		Negative	Negative	Negative	Negative	Negative	Negative
Positive Control		129±18	462±34.7	1076±45.0	854±74.9	481±83.0	568±8.4

[a] Revertants are presented as mean ± standard error from 3 plates

Expt 2 (Zeiger, et al)

Strain	Dose (µg/plate)	Revertants/Plate [a] -S9 Trial 1	-S9 Trial 2	+30% hamster S9 Trial 1	+30% hamster S9 Trial 2	+30% rat S9 Trial 1
TA100	0	159±3.5	161±11.1	151±4.7	160±10.2	170±9.0
	10		149±11.8			
	33		164±13.5			
	100	152±7.0	150±10.4	156±8.1	172±11.5	154±10.1
	333	161±12.7	154±5.4	233±15.6	225±17.5	154±3.5
	1000	154±5.8	188±4.2	335±11.9	364±21.4	157±5.8
	1666				414±32.8	
	3333	0±0.0 [b]		533±14.9	502±32.4	171±5.5
	6666	toxic		477±39.8		173±8.1
Trial Summary		Negative	Negative	Positive	Positive	Negative
Positive Control		503±5.2	1132±62.5	812±50.9	845±18.8	529±7.9
TA98	0	28±2.2	32±6.1	35±2.7		43±3.5
	10		32±4.7			
	33		41±5.5			
	100	30±3.5	32±0.3	36±3.5		46±4.5
	333	35±2.9	29±0.6	34±2.9		44±6.1
	1000	27±3.3	44±4.7	30±1.8		50±5.8
	3333	23±3.4 [b]		39±3.8		31±3.8
	6666	toxic		29±3.5		40±1.5
Trial Summary		Negative	Negative	Negative		Negative
Positive Control		677±20.6	464±26.2	770±11.3		168±3.5

[a] Revertants are presented as mean ± standard error from 3 plates

[b] Slight toxicity

Remarks: None

CONCLUSIONS

Remarks: The test substance gave a positive result only in the second experiment, which used a different concentration of S9 than the first experiment. Metabolic activation, specifically in the form of 30% Aroclor 1254-induced male Syrian hamster liver S9, was required to obtain the mutagenic response. 10% hamster S9 was ineffective, as was 10% or 30% S9 derived from the livers of pre-treated rats.

DATA QUALITY

- **Reliabilities:** 4, Not Assignable

Remarks: Secondary summary of published studies.

REFERENCES

Mortelmans, K., et al, Salmonella mutagenicity tests: II. Results from the testing of 270 chemicals. Environ. Mutagen. 8 (suppl. 7), 1-119 (1986)

Zeiger, E., et al, Salmonella mutagenicity tests: V. Results from the testing of chemicals. Environ. Mol. Mutagen. 19 (suppl. 2) 2-141, 1992

Reported in:

NTP Technical Report on the Toxicology and Carcinogenesis Studies of 2,2-bis(bromomethyl)-1,3-propanediol (FR-1138, CAS No. 3296-90-0) in F344/N Rats and B6C3F1 Mice (Feed Studies), NTP TR 452, National Toxicology Program, May 1996

OTHER

- **Last Changed:** March 26 2003
- **Order number for sorting:** 4

HUMAN HEALTH ENDPOINTS
15.5 GENETIC TOXICITY IN VITRO

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Radian Corporation
Purity: Technical grade, approximately 79% pure

METHOD

- **Method:** Galloway et al (1987)
- **Test Type:** Cytogenetic Assay (Chromosomal Aberration)
- **System of testing:** Non-bacterial
- **GLP:** No data
- **Year:** 1987
- **Species/Strain:** Chinese Hamster Ovary cells
- **Metabolic activation:** S9-mix, Rat liver cells, Aroclor induced, and cofactor
- **Concentrations tested:** 400, 500, 600, 700 µg/ml (-S9), 600, 800, 1000, 1200 µg/ml (+S9)
- **Statistical Methods:** Significance of percent cells with aberrations tested by the linear regression trend test vs. log of the dose.

Remarks:

- **Test Design**

  - **Number of replicates:** Single flask per dose, positive or equivocal results were repeated
  - **Positive Controls:** Mitomycin-C (-S9), Cyclophosphamide (+S9)
  - **Negative Control:** Solvent vehicle
  - **Solvent:** Dimethylsulfoxide
  - **Exposure Periods:** Expt 1: 18.5 hrs (-S9), 2hours (+S9); Expt 2 –S9, extra 10-12 hrs due to cell cycle delay

RESULTS

- **Result:** Positive
- **Cytotoxic concentration**

  - **With metabolic activation:** ≥ 700 µg/ml
  - **Without metabolic activation:** ≥ 1200 µg/ml

- **Genotoxic effects**

  - **With metabolic activation:** Significant ($P \leq 0.05$) dose-related increase in chromosomal aberrations.
  - **Without metabolic activation:** None

- **Statistical results:**

-S9					+S9				
Dose (µg/ml)	Total Cells	No. of Abs [a]	Abs/ Cell	Cells with Abs (%)	Dose (µg/ml)	Total Cells	No. of Abs	Abs/ Cell	Cells with Abs (%)
Harvest time: 20.5 hours [b]					Harvest time: 10.5 hours				
Summary: Negative					Summary: Positive				
Dimethylsulfoxide	100	2	0.02	2.0	Dimethylsulfoxide	100	5	0.05	5.0
Mitomycin-C (0.062 µg/ml)	50	10	0.20	16.0	Cyclophosphamide (50 µg/ml)	50	19	0.38	28.0
400	100	1	0.01	1.0	600	100	8	0.08	4.0
500	100	2	0.02	2.0	800	100	24	0.24	22.0*
600	100	0	0.00	0.0	1000	100	17	0.17	16.0*
700	0				1200	0			
				P=0.833 [c]					P≤0.001

* Positive (P<0.05)

[a] Abs = aberrations

[b] Because of significant chemical-induced cell cycle delay, incubation time prior to addition of colcemid was lengthened to provide sufficient metaphase cells at harvest

[c] Significance of percent cells with aberrations tested by the linear regression trend test vs. log of the dose

Remarks: A majority of the breaks that were observed in the study were located in the heterochromatic region of the long arm of the X chromosome. The authors did not know the reason for this preferential breakage site.

CONCLUSIONS

Remarks: The test substance induced chromosomal aberrations in CHO cells, in the presence of S9.

DATA QUALITY

- **Reliabilities:** 2, Valid with restrictions

Remarks: Study reported in literature (peer reviewed).

REFERENCES

Galloway, S.M., et al, Chromosome aberrations and sister chromatid exchanges in Chinese hamster ovary cells: Evaluations of 108 chemicals. Environ. Mol. Mutagen., 10 (suppl. 10), 1 -175, 1987

OTHER

- **Last Changed:** March 26, 2003
- **Order number for sorting:** 5

Remarks:

HUMAN HEALTH ENDPOINTS
15.6 GENETIC TOXICITY IN VITRO

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Radian Corporation
Purity: Technical grade, approximately 79% pure

METHOD

- **Method:** Galloway et al (1987)
- **Test Type:** Cytogenetic Assay (Sister Chromatid exchange Test)
- **System of testing:** Non-bacterial
- **GLP:** No data
- **Year:** 1987
- **Species/Strain:** Chinese Hamster Ovary cells
- **Metabolic activation:** S9-mix, Rat liver cells, Aroclor induced, and cofactor
- **Concentrations tested:** 16.7, 50, 167, 500 µg/ml (-S9), 800, 1000, 1200 µg/ml (+S9)
- **Statistical Methods:** Significance of relative SCE/chromosome tested by the linear regression trend test vs. log of the dose.

Remarks:

- **Test Design**

  - **Number of replicates:** Single flask per dose, positive or equivocal results were repeated
  - **Positive Controls:** Mitomycin-C (-S9), Cyclophosphamide (+S9)
  - **Negative Control:** Solvent vehicle
  - **Solvent:** Dimethylsulfoxide
  - **Exposure Periods:** Expt 1: 26.3 hrs (-S9), 2hours (+S9); Expt 2 –S9, incubation time increased at 167 and 500 µg/kg due to significant chemical-induced cell cycle delay.

RESULTS

- **Result:** Equivocal
- **Cytotoxic concentration**

  - **With metabolic activation:** 1200 µg/ml
  - **Without metabolic activation:** ≥ 500 µg/ml

- **Genotoxic effects**

  - **With metabolic activation:** None
  - **Without metabolic activation:** Very slight increases in SCEs occurred at toxic levels

- **Statistical results:**

Dose (µg/ml)	Total Cells	No. of Chromosomes	No. of SCEs [a]	SCEs/ Chromosome	SCEs/ Cell	Hrs in BrdU [a]	Relative change of SCEs/Chromosome [b] (%)
-S9							
Summary: Negative							
Solvent	50	1038	496	0.47	9.9	26.3	
Positive control (0.005 µg/ml)	25	519	692	1.33	27.7	26.3	179.03
16.7	50	1041	485	0.46	9.7	26.3	-2.50
50	50	1042	498	0.47	10.0	26.3	0.02
167	50	1050	545	0.51	10.9	33.5 [c]	8.62
500	0					33.5 [c]	
				P=0.077 [d]			
+S9							
Summary: Equivocal							
Solvent	50	1050	496	0.47	9.9	25.5	
Positive control (1.5 µg/ml)	25	523	840	1.6	33.6	25.5	240.00
800	50	1048	556	0.53	11.1	25.5	12.31
1000	50	1047	590	0.56	11.8	25.5	19.29
1200 [e]	50	1046	574	0.54	11.5	25.5	16.17
				P=0.004			

[a] SCE = sister chromatid exchange, BrdU = bromodeoxyuridine

[b] SCEs/chromosome in treated cells versus SCEs/chromosome in solvent control cells

[c] Due to chemical induced cell cycle delay, incubation time was extended to provide sufficient cells for scoring

[d] Significance of relative SCEs/chromosome tested by the linear regression trend test vs. log of the dose

[e] Marked toxicity noted at this dose

Remarks: None

CONCLUSIONS

Remarks: The test substance did not induce sister chromatid exchanges in CHO cells without S9, even at doses that induced toxicity and marked cell cycle delay. Very slight increases in SCEs occurred at toxic levels with S9. The top dose, 1.2 mg/ml, reduced confluence by about 75%.

DATA QUALITY

- **Reliabilities:** 2, Valid with restrictions

Remarks: Study reported in literature (peer reviewed).

REFERENCES

Galloway, S.M., et al, Chromosome aberrations and sister chromatid exchanges in Chinese hamster ovary cells: Evaluations of 108 chemicals. Environ. Mol. Mutagen., 10 (suppl. 10), 1 -175, 1987

OTHER

- **Last Changed:** March 26, 2003
- **Order number for sorting:** 6

Remarks:

HUMAN HEALTH ENDPOINTS
15.7 GENETIC TOXICITY IN VITRO

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Dead Sea Bromine Co.

METHOD

- **Method:**
- **Test Type:** Reverse Mutation Assay (Ames Test)
- **System of testing:** Bacterial
- **GLP:** No
- **Year:** 1977
- **Species/Strain:** Salmonella typhimurium strains TA98, TA100, TA1535 and TA1537
- **Metabolic activation:** S9-mix, Rat liver cells, Aroclor induced
- **Concentrations tested:** 1, 10, 100, 500 µg/plate
- **Statistical Methods:** No data

Remarks:

- **Test Design**

  - **Number of replicates:** 2
  - **Positive Controls:** N-methyl-N'-nitroso-N-nitrosoguanidine (-S9)
    Daunomycin (-S9)
    Acridine-orange (+S9)
  - **Negative Control:** Solvent vehicle

- **Solvent:** Dimethylsulfoxide

RESULTS

- **Result:** Negative
- **Cytotoxic concentration**

  - **With metabolic activation:** > 500 µg/plate
  - **Without metabolic activation:** > 500 µg/plate

- **Genotoxic effects**
  - **With metabolic activation:** None
  - **Without metabolic activation:** None

- **Statistical results:** All strains yielded colony counts that were not significantly different from the comparable control counts.

Type of experiment	Microsome preparation	Compound*	µg/plate	No. of revertant colonies per plate**			
				TA98	TA100	TA1535	TA1537
Solvent controls	-	DMSO		30; 58	203; 247	91; 103	14; 16
	+	DMSO		32; 55	185; 164	19; 21	8; 12
Positive controls	-	MNNG	2		10038; 12500	7500; 8500	
	-	Daunomycin	5	474; 510			
	-	AO	20	96; 107			
	+	AO	20	1447; 1710	1778; 1965	28; 42	95; 103
Test runs	-	DBNPG	500	34; 48	222; 248	89; 90	19; 19
	+	''	500	35; 49	111; 154	14; 24	7; 9
	-	''	100	42; 45	238; 284	62; 93	6; 19
	+	''	100	45; 56	167; 183	20; 21	8; 9
	-	''	10	37; 39	212; 241	52; 74	25; 26
	+	''	10	52; 56	164; 168	17; 22	7; 13
	-	''	1	41; 51	219; 233	56; 67	9; 11
	+	''	1	49; 52	175; 185	19; 23	7; 11

MMNG = N-methyl-N'-nitroso-N-nitrosoguanidine, AO = Acridine-orange

Remarks: None

CONCLUSIONS

Remarks: None

DATA QUALITY

- **Reliabilities:** 3, Not reliable

Remarks: No details on purity of test substance or lot number used. Test not subject to QA overview. Substance only tested to 500 µg/plate.

REFERENCES

Israel Institute for Biological Research, Ames salmonella/microsome mutagenicity assay for Dibromo neo-pentyl glycol (DBNPG), October 1977

OTHER

- **Last Changed:** October 3 2002
- **Order number for sorting:** 7

Remarks:

HUMAN HEALTH ENDPOINTS
15.8 GENETIC TOXICITY IN VITRO

TEST SUBSTANCE

- Monobromopentaerythritol

Remarks: Purity: 98.2%, Lot No.: 564-347-01
Source: Bromine Compounds Ltd.
This study is provided as additional information from a structurally similar compound.

METHOD

- **Method:** OECD 471, 92/69/EEC Method B14, US EPA, Method: HG-Gene Muta– S. typhimurium: The Salmonella typhimurium reverse mutation assay, 1984
- **Test Type:** Reverse Mutation Assay
- **System of testing:** Bacterial
- **GLP:** Yes
- **Year:** 1996
- **Species/Strain:** Salmonella typhimurium strains TA98, TA100, TA1535 and TA1537
- **Metabolic activation:** S9-mix, Rat liver cells, Aroclor induced
  S9-mix, Hamster liver cells, uninduced
- **Concentrations tested:** 50, 150, 500, 1500, 5000 µg/plate
- **Statistical Methods:** None

Remarks:

- **Test Design**
  - **Number of replicates:** 3
  - **Positive Controls:** N-ethyl-N'-nitroso-N-nitrosoguanidine (TA100, TA1535, -S9)
    9-aminoacridine (TA1537, -S9)
    2-nitrofluorene (TA98, -S9)
    2-aminoanthracene (TA98, TA100, TA 1535, TA1537, +S9)
    Congo red (TA98, +S9)
  - **Negative Control:** Solvent vehicle
- **Solvent:** Dimethylsulfoxide

RESULTS

- **Result:** Negative
- **Cytotoxic concentration**
  - **With metabolic activation:** > 5000 µg/plate
  - **Without metabolic activation:** > 5000 µg/plate
- **Genotoxic effects**
  - **With metabolic activation:** None
  - **Without metabolic activation:** None

- **Statistical results:**

Strain	Dose level (µg/plate)	Revertants/Plate [a]							
		Test 1				Test 2			
		-S9	+10% Rat S9	+10% Hamster S9	+30% Hamster S9	-S9	+10% Rat S9	+10% Hamster S9	+30% Hamster S9
TA1535	5000	12±1.2	14±4.2	9±3.2	15±0.0	14±4.2	18±3.6	16±6.7	13±2.6
	1500	14±2.1	13±1.2	12±1.5	11±3.2	13±4.5	15±4.2	15±3.0	15±2.9
	500	9±3.8	12±2.3	11±3.1	10±4.7	9±6.8	12±2.9	16±2.5	16±3.8
	150	9±2.3	12±3.1	12±5.3	10±5.5	10±0.6	13±1.7	13±2.0	15±2.1
	50	8±1.2	12±2.6	13±1.7	11±2.6	11±1.0	13±3.5	17±4.5	11±1.5
	0	11±3.1	16±0.0	15±6.1	16±1.5	14±5.5	16±0.6	17±0.6	15±2.5
	Solvent	12±2.0	12±1.5	12±0.6	16±2.1	13±2.3	17±1.5	12±2.5	18±2.1
	ENNG (5.0)	516±71.7				1151±130.3			
	AA (2.0)		92±8.0	102±60.1	131±9.3		120±19.7	86±17.6	146±36.0
TA1537	5000	12±6.8	13±1.5	10±3.0	8±3.1	9±1.0	15±2.6	11±1.7	8±5.6
	1500	13±3.0	12±2.3	9±2.5	12±4.5	10±1.5	14±6.7	11±1.2	12±1.5
	500	9±1.2	7±1.0	11±4.2	11±2.6	8±1.5	11±0.6	10±2.6	11±3.8
	150	12±0.6	11±1.2	10±2.3	11±1.7	11±6.5	12±7.0	14±4.4	12±5.0
	50	10±1.5	9±3.2	12±1.5	9±2.1	7±2.1	11±5.6	12±2.6	11±2.5
	0	9±1.0	14±4.0	9±1.0	11±4.4	14±3.6	15±5.5	12±3.2	15±4.0
	Solvent	11±2.5	11±1.7	13±3.6	12±3.2	13±3.5	15±4.0	17±0.6	17±3.1
	9 AC (80.8)	X				X			
	AA (2.0)		45±4.4	151±41.6	99±43.9		72±18.7	135±14.4	98±23.4
TA98	5000	32±4.0	27±4.6	28±2.0	25±3.1	24±4.9	28±4.6	22±2.1	27±7.2
	1500	22±6.5	23±3.8	26±1.7	27±9.0	25±4.0	25±2.3	22±3.2	24±2.6
	500	26±7.5	21±4.2	22±3.5	25±4.0	22±5.5	24±6.4	21±3.2	28±1.7
	150	18±0.6	26±6.6	26±0.6	26±4.6	23±2.5	26±5.2	23±3.2	24±4.5
	50	23±4.5	25±4.7	20±5.5	22±3.2	19±2.1	26±3.0	21±1.7	25±5.3
	0	22±2.6	22±2.1	28±4.5	23±4.7	23±4.0	23±2.9	24±4.0	25±2.5
	Solvent	25±3.8	24±3.2	27±4.0	27±4.4	24±2.1	27±4.2	25±2.0	27±3.8
	NF (1.0)	152±2.1				388±15.6			
	AA (0.5)		159±25.3	367±32.7	114±0.0		175±8.1	675±15.3	430±71.2
	CR (200.0)		24±3.0	43±3.2	124±7.0		48±6.7	57±2.1	126±44.7
TA100	5000	107±2.6	113±6.8	128±5.1	117±7.8	118±11.8	111±0.6	117±10.1	146±9.8
	1500	104±4.0	114±27.7	117±25.0	118±6.1	110±9.2	110±16.3	124±6.2	127±2.1
	500	112±14.6	116±4.4	114±7.6	117±10.7	106±6.0	113±11.9	109±6.9	122±27.3
	150	114±6.8	108±5.5	117±6.6	119±2.3	114±7.5	124±12.1	119±2.0	134±6.8
	50	110±6.7	113±10.2	105±3.2	105±8.1	113±6.1	111±1.5	110±9.6	127±10.5
	0	109±5.0	103±12.2	129±20.8	119±8.5	127±6.1	128±7.5	133±12.6	136±9.3
	Solvent	110±11.7	129±4.2	109±11.2	130±7.5	112±6.7	120±4.0	129±10.3	142±8.9
	ENNG (3.0)	328±4.0				596±31.3			
	AA (1.0)		332±28.1	560±135.2	293±107.7		433±56.1	915±124.4	801±130.8

[a] Data presented as mean ± standard deviation

ENNG N-ethyl-N'-nitro-N-nitrosoguanidine, 9 AC 9-aminoacridine, NF 2-nitrofluorene, AA 2-aminoanthracene, CR Congo red

X Too many colonies to count accurately

Remarks: No substantial increases in revertant colony numbers of any of the tester strains were observed following treatment with monobromopentaerythritol at any dose level, in the presence or absence of S-9 mix, in either mutation test. The concurrent positive control compounds demonstrated the sensitivity of the assay and the metabolizing activity of the liver preparations.

CONCLUSIONS

Remarks: It is concluded that, when tested in dimethylsulfoxide, the test substance shows no evidence of mutagenic activity in this bacterial system.

DATA QUALITY

- **Reliabilities:** 1, Reliable without restrictions

Remarks: Study conducted under GLP to OECD/EC/EPA test guidelines by Huntingdon Life Sciences.

REFERENCES

Huntingdon Life Sciences Ltd., Monobromopentaerythritol, Bacterial Mutation Assay, January 1996

OTHER

- **Last Changed:** October 3 2002
- **Order number for sorting:** 8

Remarks: S-9 is usually only prepared from a rat previously treated with a compound (Aroclor) known to induce a high level of enzymic activity. However in this study S-9 mix using liver fraction from uninduced Syrian hamsters was also used as it was suspected that the test substance may be metabolised to a mutagen by hamster S -9 but not by rat.

HUMAN HEALTH ENDPOINTS
16.1 REPEATED DOSE TOXICITY

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Dow Chemical Company, Lot No.: 840429-162, Purity: approximately 78.6%, Identity of impurities: 2,2-bis (hydroxymethyl)-1-bromo-3-hydroxypropane (6.6%), 2,2-bis (bromomethyl)-1-bromo-3-hydroxypropane (6.9%), pentaerythritol (0.2%), dimers and structural isomers (7.7%).

METHOD

- **Method/guideline followed:** Similar to OECD 453, however neither neurological nor ocular assessments were made during the study.
- **Test type:** 2 year repeated dose toxicity and carcinogenicity study
- **GLP:** Yes
- **Year:** 1991
- **Species:** Rat, Mouse
- **Strain:** F344/N (rat), B6C3F_1 (Mouse)
- **Route of administration:** Oral feed
- **Duration of test:** 2 years
- **Doses/concentration levels:**

Rat – 0, 2500, 5000, 10000 ppm
Rat (stop exposure evaluation) – 20000 ppm
Mouse – 0, 312, 625, 1250 ppm

- **Sex:** Male/female (rat/mouse), male rats used in stop-exposure evaluation
- **Exposure period:** 104/105 weeks, 3 months for stop-exposure evaluation
- **Frequency of treatment:** Continuous, ad libitum
- **Control group and treatment:**

Rats - 70 male/60 female, feed control
Mice – 60 male/60 female, feed control

- **Post exposure observation period:** None for 2-year exposure rats and mice, 21 months for stop-exposure rats

- **Statistical methods:** Kaplan & Meier product-limit procedure (probability of survival, estimate of neoplasm incidence). Cox's method for testing 2 groups for equality and Tarone's life table test (dose-related trends). Dinse & Lagakos, Dinse & Haseman prevalence analysis (time-specific neoplasm incidences). Fisher exact test, Cochran-Armitage trend test. Dunnett and Williams parametric multiple comparisons (organ & body weights). Shirley and Dun's non-parametric multiple comparison methods (clinical chemistry, urinalysis, spermatid and epididymal spermatozoa data). Jonckheere's test (significance of dose-related trends). Mann-Whitney U test (average severity values). Morrison multivariate analysis of variance (treatment effects).

Remarks:

- **Test subjects**

  - **Age at study initiation:** Rats/mice – 6 weeks old

  - **No. of animals/sex/dose:** Rats/mice - 60 male/60 female per dose. 70 male rats were used in the stop exposure evaluation.

- **Study Design**

  - **Vehicle:** Feed

  - **Satellite groups and reasons they were added:** A group of 70 male rats received 20000 ppm in feed for 3 months. 10 rats (and 10 male controls) were evaluated at this time and the remaining rats were fed control feed for the remainder of the study. Group added to evaluate the potential for progression or regression of urinary bladder and kidney lesions.

  - **Clinical observations performed and frequency:** Observed twice daily; body weights and clinical observations recorded initially, weekly for weeks 2 to 13, monthly thereafter, and at the end of the studies. Feed consumption was measured every 4 weeks by cage.

  - **Organs examined at necropsy (macroscopic and microscopic):** Complete histopathologic examinations were performed on all animals necropsied. In addition to gross lesions, tissue masses and associated lymph nodes, the tissues examined included: adrenal gland, bone and marrow, brain, esophagus, gallbladder (mice), heart, kidney, large intestine (cecum, colon and rectum), liver, lung, lymph nodes (mandibular or mesenteric), mammary gland, nose, ovary, pancreas, parathyroid gland, pituitary gland, preputial or clitoral gland, prostate gland, salivary gland, skin, small intestine, (duodenum, jejunum and ileum), spleen, stomach (forestomach and glandular), testis with epididymis and seminal vesicle, thymus, thyroid gland, trachea, urinary bladder and uterus.

RESULTS

- **NOAEL (rats):** Not achieved
- **LOAEL (rats):** 2500 ppm
- **NOAEL (mice):** Not achieved
- **LOAEL (mice):** 312 ppm

- **Actual dose received by sex:**
  Rats – 100, 200, 430 mg/kg bw (male); 115, 230, 460 mg/kg (female); 800 mg/kg bw (Stop exposure males)
  Mice – 35, 70, 140 mg/kg bw (male); 40, 80, 170 mg/kg bw (female)

- Toxic response/effects by dose level:

- Statistical results, as appropriate:

Remarks:

- **Body weight:**
  Rats – Mean body weights of exposed male and female rats receiving 10000 ppm and stop exposure males receiving 20000 were lower than those of the controls throughout most of the study.
  Mice – Mean body weights of exposed male and female mice were similar to controls throughout the study. Final mean bodyweights were also generally similar to those of controls.

- **Food/water consumption:**
  Rats – In the continuous exposure study, feed consumption by exposed rats was generally similar to that by the controls throughout the study. In 20000-ppm stop exposure males, the feed consumption was lower than that by controls.
  Mice – Feed consumption by exposed male and female mice was similar to that by controls.

- **Description, severity, time of onset and duration of clinical signs:**
  Rats – Clinical findings included skin and subcutaneous tissue masses on the face, tail and the ventral and dorsal surfaces of exposed rats.
  Mice – Clinical findings included swelling, discharge and tissue masses involving the eye in exposed mice.

- **Ophthalmologic findings, incidence and severity:** No data

- **Hematologic findings, incidence and severity:** No data

- **Clinical biochemistry findings, incidence and severity:** No data

- **Mortality and time to death:**
  Rats – Survival of 5000 and 1000 continuous exposure males and females and 20000-ppm stop exposure males was significantly lower than controls.
  Mice – Survival of 1250-ppm males and females was significantly lower than controls.

Rat survival

	Males					Females			
	Dose (ppm)					Dose (ppm)			
	0	2500	5000	10000	20000[a]	0	2500	5000	10000
Animals initially in study	70	60	60	60	70	60	60	60	60
3 month interim evaluation[b]	10	0	0	0	10				
15 month interim evaluation[b]	9	7	9	5	0	10	9	7	8
Moribund	24	30	36	43	55	14	22	27	41
Natural deaths	1	3	2	11	5	0	2	3	6
Animals surviving to study termination	26	20	13	1	0	36	27	23	5
Percentage probability of survival at the end of the study[c]	51	38	26	2	0	72	53	43	10
Mean survival (days)[d]	688	652	669	587	544	711	701	676	630

Survival analysis[e]	P<0.001	P=0.126	P=0.010	P<0.001	P<0.001	P<0.001	P=0.095	P=0.005	P<0.001

[a] Ten male rats receiving 20000 ppm test substance in feed were evaluated at 3 months. The remaining 60 male rats received control feed until the end of the 2 year study. [b] censored from survival analysis. [c] Kaplan-Meier determinations. [d] Mean of all deaths (uncensored, censored and terminal sacrifice). [e] The result of the life table trend test (Tarone, 1975) is in the control column and the results of the life table pairwise comparisons (Cox, 1972) with the controls are in the dosed columns.

Mouse survival

	Males				Females			
	Dose (ppm)				Dose (ppm)			
	0	312	625	1250	0	312	625	1250
Animals initially in study	60	60	60	60	60	60	60	60
15 month interim evaluation [a]	10	9	10	10	8	10	9	10
Accidental deaths [a]	0	0	0	1	0	0	0	0
Missing [a]	0	0	0	1	0	0	0	0
Moribund	3	12	11	13	9	14	14	29
Natural deaths	5	3	4	5	6	6	11	10
Animals surviving to study termination	42	36	35	30	37	30	26	11
Percentage probability of survival at the end of the study[b]	84	71	70	63	71	60	51	22
Mean survival (days) [c]	710	675	698	684	690	685	691	625
Survival analysis[d]	P=0.054	P=0.169	P=0.174	P=0.038	P<0.001	P=0.422	P=0.117	P<0.001

[a] Censored from survival analyses. [b] Kaplan-Meier determinations. [c] Mean of all deaths (uncensored, censored and terminal sacrifice). [d] The result of the life table trend test (Tarone, 1975) is in the control column and the results of the life table pairwise comparisons (Cox, 1972) with the controls are in the dosed columns.

- **Gross pathology incidence and severity:** No data, although it is likely that some or all the neoplasms found would be visible macroscopically.

- **Organ weight changes:**
  Rats – At the 3-month interim evaluation, the right kidney and liver weights of 20000-ppm stop-exposure males were significantly lower (P ≤0.01) than controls. The right kidney and liver weights relative to bodyweight in this group were significantly higher (P≤0.01) than controls. At the 15-month interim evaluation, the relative right kidney weights in 10000-ppm males and females were significantly higher than controls. The relative liver weights in 5000 and 10000-ppm males and 2500, 5000 and 10000 females were significantly higher (P≤0.05 & P≤0.01, respectively) than controls.
  Mice – No significant differences.

- **Histopathology incidence and severity:**

  Rats:
  Skin: The incidence of squamous cell papilloma in 20000-ppm stop-exposure males was significantly greater than controls. The incidences of keratocanthoma and of squamous and basal cell neoplasms (combined) in 5000 and 10000-ppm continuous-exposure males and in the 20000-ppm stop-exposure males were significantly greater than controls.

  Mammary Gland: The incidences of benign mammary gland neoplasms (fibroadenoma and fibroadenoma or adenoma (combined)) were significantly greater in the stop-exposure group of male rats and in all continuous-exposure groups of male rats than controls. In female rats, the incidences of fibroadenoma and of fibroadenoma, adenoma, or carcinoma (combined) in all exposed groups

were greater than controls. The incidences of multiple fibroadenoma in all exposed female groups were greater than controls.

Zymbal's Gland: The incidences of Zymbal's gland adenoma in 10000-ppm males and of adenoma or carcinoma in 20000-ppm stop-exposure males were significantly greater than those in the controls.

Oral cavity (Pharynx, Tongue and Gingiva), Esophagus and Forestomach: The incidences of squamous cell papilloma of the oral cavity in exposed males (continuous and stop-exposure) were significantly greater than controls. Additionally, the incidences of squamous cell papilloma of the esophagus in male and female rats exposed to 10000 ppm were significantly greater than controls. The incidences of squamous cell neoplasms of the oral cavity in 20000-ppm stop-exposure males were similar to those in 10000-ppm males. Squamous cell papilloma of the forestomach occurred in exposed groups of rats, but the incidence was only significant in 20000-ppm stop-exposure males.

Small and Large Intestine: There was a significant positive trend in the incidences of adenoma (adenomateous polyp) in the large intestine of males. In 20000-ppm stop-exposure males, the incidence of adenoma and adenoma or carcinoma (combined) of the large intestine was significantly greater than controls.

Kidney and Urinary Bladder: In male rats, the incidence of renal tubule adenoma in the 10000-ppm group was marginally but significantly greater than controls.

Lung: The incidences of alveolar/bronchiolar adenoma or carcinoma (combined) in 10000-ppm continuous exposure males and 20000-ppm stop-exposure males were significantly greater than controls.

Thyroid Gland: In males, the incidence of follicular cell adenoma in the 20000-ppm stop-exposure group and the incidence of follicular cell adenoma or carcinoma in 5000-ppm continuous-exposure males, 20000-ppm stop-exposure males and 10000-ppm females were significantly greater than controls.

Hematopoietic System: The incidences of mononuclear cell leukemia in male rats from the 5000 and 10000-ppm continuous-exposure groups and the 20000 stop-exposure group were significantly greater than controls.

Pancreas: The incidences of hyperplasia in all exposed groups of male rats were significantly greater than controls.

Summary of study findings - Rats

	Males Dose (ppm)					Females Dose (ppm)			
	0	2500	5000	10000	20000	0	2500	5000	10000
Nonneoplastic effects									
Kidney									
Focal atrophy	0/51	0/53	0/51	5/55	0/59	0/50	2/51	1/53	7/52
Papillary degeneration	0/51	5/53	30/51	29/55	16/59	0/50	1/51	3/53	17/52
Papillary epithelial hyperplasia	10/51	20/53	25/51	47/55	21/59	0/50	1/51	1/53	7/52
Pelvis, transitional epithelium, hyperplasia	0/51	0/53	0/51	4/55	4/59				
Lung									
Alveolar/bronchiolar hyperplasia	3/51	4/53	5/51	7/55	14/60				
Thyroid Gland									
Follicular cell hyperplasia	1/51	0/53	2/51	5/55	6/59				
Seminal Vesicle									
Hyperplasia	1/51	6/53	4/51	16/55	33/60				
Pancreas									
Focal hyperplasia	3/51	9/53	12/51	14/53	27/59				
Forestomach									
Mucosal hyperplasia	4/51	12/53	6/51	6/55	6/59				
Urinary Bladder									
Hyperplasia	0/51	0/53	1/51	3/55	10/59				
Neoplastic effects									
Skin									
Squamous cell papilloma, keratocanthoma, trichoepithelioma, basal cell adenoma, basal or squamous cell carcinoma	4/51	6/53	14/51	24/55	21/60				
Skin, Subcutaneous Tissue									
Fibroma, fibrosarcoma or sarcoma	2/51	9/53	13/51	16/55	10/60				
Mammary Gland									
Fibroadenoma or adenoma	0/51	4/53	7/51	7/55	5/60				
Fibroadenoma						25/50	45/51	46/53	45/52
Zymbal's Gland									
Adenoma or carcinoma	2/51	1/53	4/51	5/55	15/60				
Oral Cavity (pharynx, tongue or gingiva)									
Squamous cell papilloma or carcinoma	0/51	4/53	9/51	10/55	13/60	2/50	3/51	5/53	6/52
Esophagus									
Squamous cell papilloma	0/51	0/53	1/51	5/55	0/60	0/50	0/51	1/53	10/52
Forestomach									
Squamous cell papilloma	0/51	0/53	0/51	1/55	5/60				
Large Intestine									
Adenoma or carcinoma	0/51	0/53	3/51	4/55	11/59				
Small Intestine									
Adenoma or carcinoma	0/51	0/53	0/51	2/53	5/59				
Malignant Mesothelioma	0/51	3/53	8/51	9/55	25/60				
Urinary Bladder									
Transitional cell papilloma or carcinoma	0/51	0/53	1/51	3/55	2/59				
Lung									
Alveolar/bronchiolar adenoma or carcinoma	1/51	1/53	3/51	4/55	7/60				
Squamous cell carcinoma	0/51	0/53	0/51	0/55	3/60				
Thyroid Gland									
Follicular cell adenoma or carcinoma	0/51	2/53	6/51	3/55	9/59	0/50	0/51	2/53	4/52
Seminal Vesicle									
Adenoma or carcinoma	0/51	0/53	0/51	0/55	2/60				
Hematopoietic System									
Mononuclear cell leukemia	27/51	29/53	40/51	34/55	25/60				
Uncertain effects									
Kidney (renal tubule)									
Adenoma	0/51	0/53	1/51	3/55	1/59				
Pancreas									
Acinar cell adenoma	1/51	2/53	4/51	3/53	3/59				

Mice:
Harderian Gland: The incidences of harderian gland adenoma in male and female mice exposed to 625 and 1250 ppm were significantly greater than controls. The incidence of harderian gland carcinoma in 1250-ppm females was significantly greater that controls. In 625-ppm and 1250-ppm males and in all female exposure groups, the incidences of adenoma or carcinoma 9combined) were significantly greater than those in the control groups.

Lung: The incidences of alveolar/bronchiolar adenoma and of alveolar/bronchiolar adenoma of carcinoma (combined) in 1250-ppm males and females and 625 ppm females were significantly greater than controls. In males exposed to 1250 ppm the incidences of multiple adenoma and of alveolar/bronchiolar carcinoma were significantly greater than controls. At 2 years, the incidences of alveolar epithelial hyperplasia in 625 and 1250-ppm females were significantly greater than controls.

Skin: The incidence of subcutaneous tissue sarcoma and the combined incidences of fibrosarcoma or sarcoma in 1250-ppm female mice were significantly greater than controls.

Forestomach: The incidences of squamous cell papilloma of the forestomach in 625 and 1250-ppm female mice were significantly greater that controls. In 1250-ppm males, the incidence of squamous cell papilloma or squamous cell carcinoma (combined) was significantly greater than controls.

Other: The incidence of hemangioma or hemangiosarcoma (combined) was significantly increased in 1250-ppm female mice.

Summary of study findings - Mice

	Males Dose (ppm)				Females Dose (ppm)			
	0	312	625	1250	0	312	625	1250
Nonneoplastic effects								
Lung								
Alveolar epithelium, hyperplasia					1/52	3/50	8/51	15/50
Neoplastic effects								
Harderian Gland								
Adenoma or carcinoma	4/50	7/51	16/50	22/49	3/52	12/50	13/51	19/50
Skin, Subcutaneous Tissue								
Sarcoma					0/52	1/50	4/51	11/50
Lung								
Alveolar/bronchiolar adenoma or carcinoma	15/50	11/51	16/50	25/49	5/52	5/50	15/51	19/50
Kidney (renal tubule)								
Adenoma	0/50	0/51	3/50	2/49				
Uncertain effects								
Forestomach								
Squamous cell papilloma or carcinoma	0/50	3/51	3/50	4/49				
Squamous cell papilloma					0/52	1/50	5/51	3/50
Mammary Gland								
Carcinoma					0/52	0/50	1/51	3/50
Circulatory System								
Hemangioma and hemangiosarcoma					1/52	2/50	0/51	5/50

CONCLUSIONS

Remarks:

There was clear evidence of carcinogenic activity of the test substance in male F344/N rats based on increased incidences of neoplasms of the skin, subcutaneous tissue, mammary gland, Zymbal's gland, oral cavity, esophagus, forestomach, small and large intestines, mesothelium, urinary bladder, lung, thyroid gland and seminal vesicle, and the increased incidence of mononuclear cell leukemia.

There was clear evidence of carcinogenic activity of the substance in female F344/N rats based on increased incidences of neoplasms of the oral cavity, esophagus, mammary gland and thyroid gland.

There was clear evidence of carcinogenic activity of the substance in male B6C3F_1 mice based on increased incidences of neoplasms of the harderian gland, lung and kidney.

There was clear evidence of carcinogenic activity of the substance in female B6C3F_1 mice based on increased incidences of neoplasms of the harderian gland, lung and subcutaneous tissue.

Slight increases in the incidences of neoplasms in the pancreas and kidney in male mice and the forestomach, mammary gland and circulatory system in female mice may also have been related to treatment.

DATA QUALITY

- 1, Valid without restrictions

Remarks: Well-conducted study performed by Southern Research Institute.

REFERENCES:

NTP Technical Report on the Toxicology and Carcinogenesis Studies of 2,2-bis(bromomethyl)-1,3-propanediol (FR-1138, CAS No. 3296-90-0) in F344/N Rats and B6C3F1 Mice (Feed Studies), NTP TR 452, National Toxicology Program, May 1996

OTHER:

- **Last Changed:** March 26, 2003
- **Order number for sorting:** 1

Remarks:

HUMAN HEALTH ENDPOINTS
16.2 REPEATED DOSE TOXICITY

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Dow Chemical Company, Lot No.: 840429-162, Purity: approximately 78.6%, Identity of impurities: 2,2-bis (hydroxymethyl)-1-bromo-3-hydroxypropane (6.6%), 2,2-bis (bromomethyl)-1-bromo-3-hydroxypropane (6.9%), pentaerythritol (0.2%), dimers and structural isomers (7.7%).

METHOD

- **Method/guideline followed:** Similar to OECD 408, however ophthalmologic examination, sensory reactivity, grip strength and motor activity assessments were not performed.
- **Test type:** 13 Week repeated dose oral toxicity study
- **GLP:** Yes
- **Year:** 1986
- **Species:** Rat, Mouse
- **Strain:** F344/N (rat), B6C3F_1 (Mouse)
- **Route of administration:** Oral feed
- **Duration of test:** 13 weeks
- **Doses/concentration levels:**

Rat – 0, 1250, 2500, 5000, 10000, 20000 ppm
Mouse – 0, 625, 1250, 2500, 5000, 10000 ppm

- **Sex:** Male/female (rat/mouse)
- **Exposure period:** 13 weeks
- **Frequency of treatment:** Continuous, ad libitum
- **Control group and treatment:** 10 male/10 female, feed control (rat/mouse)
- **Post exposure observation period:** None
- **Statistical methods:** Williams' or Shirley's test, Dunnett's or Dunn's test (body weight, organ weights, clinical chemistry, urinalysis, sperm data), Fisher exact test (incidence of lesions), Mann Whitney U test (severity values), multivariate analysis of variables (vaginal cytology data)

Remarks:

- **Test subjects**

  - **Age at study initiation:** Rat – 6 weeks
    Mouse – 6 weeks

  - **No. of animals/sex/dose:** 10 male/10 female per dose (rat/mouse)

- **Study Design**

  - **Vehicle:** Feed

  - **Satellite groups and reasons they were added:** Special clinical chemistry and Urinalysis study group (rats only)

  - **Clinical observations performed and frequency:** Observed twice daily; animals were weighed initially, weekly and at the end of the studies. Clinical observations were recorded weekly. Feed consumption was measured weekly by cage. At the end of the 13-week studies, blood was collected from the retro-orbital sinus and urine was collected from all rats and mice. In the special study rats, blood was collected on days 3, 15, 30, 60 and at study termination. Urine samples were collected on days 3, 15, 30 and 60 and at study termination. Additional urine samples were collected for measurement of urinary concentrating ability following 16-hour water deprivation periods.
    Clinical chemistry: albumin, albumin/globulin ratio, creatinine (rats only), globulin, glucose, total protein, and urea nitrogen.
    Urinalysis: glucose, protein, specific gravity and volume.

    Sperm and vaginal fluid samples were evaluated in 0, 5000, 10000 and 20000-ppm rats and 0, 2500, 5000 and 10000-ppm mice at the end of the studies. The parameters evaluated in males were sperm count, morphology and motility. The right cauda, right epididymis and right testis were weighed. Vaginal fluid samples were collected for up to 7 consecutive days prior to the end of the studies for vaginal cytology evaluations. The parameters evaluated in females were relative frequency of estrous stages and estrous cycle length.

  - Organs examined at necropsy (macroscopic and microscopic): Organs weighed included brain, heart right kidney, liver lung, spleen, right testis and thymus. Complete histopathologic examinations were performed on all control rats and mice, 20000-ppm rats and 10000-ppm mice. In addition to gross lesions, tissue masses and associated lymph nodes, the tissues examined included: adrenal gland, bone and marrow, brain, oesophagus, gallbladder (mice), heart, kidney, large intestine (cecum, colon and rectum), liver, lung, lymph nodes (mandibular or mesenteric), mammary gland, nose, ovary, pancreas, parathyroid gland, pituitary gland, preputial or clitoral gland (rats), prostate gland, salivary gland, skin, small intestine, (duodenum, jejunum and ileum), spleen, stomach (forestomach and glandular), testis with epididymis and seminal vesicle, thymus, thyroid gland, trachea, urinary bladder and uterus. The kidney and urinary bladder of all other rats and mice were also examined.

RESULTS

- **NOEL (rats):** 1250 ppm
- **LOEL (rats):** 2500 ppm
- **NOEL (mice):** 625 ppm (males), not achieved for females
- **LOEL (mice):** 1250 ppm (males), 625 ppm (females)

- **Actual dose received by sex:**
  Rats - 100, 200, 400, 800, 1700 mg/kg bw (male), 100, 200, 400, 800, 1630 mg/kg (female)
  Mice – 100, 200, 500, 1300, 3000 mg/kg bw (male), 140, 300, 600, 1200, 2900 mg/kg bw (female)

- Toxic response/effects by dose level:

- Statistical results, as appropriate:

Remarks:

- **Body weight:**
  Rats – The final mean body weights and weight gains of 5000, 10000 and 20000-ppm males and females were significantly lower than those of the controls.
  Mice – The final mean body weights and body weight gains of males and females receiving 1250, 2500, 5000 or 10000-ppm and of females receiving 625 ppm were significantly lower than those of the controls.

- **Food/water consumption:**
  Rats – Feed consumption by exposed animals was lower than that by controls at week 1, but was generally similar to or slightly higher than that of controls at week 13.
  Mice – Feed consumption by exposed mice was generally higher than that by controls throughout the study.

- **Description, severity, time of onset and duration of clinical signs:**
  Rats – No chemical-related clinical findings were observed.
  Mice – Clinical findings included abnormal posture and hypoactivity in 10000-ppm male and female mice.

- **Ophthalmologic findings, incidence and severity:** No data

- **Hematologic findings, incidence and severity:** No data

- **Clinical biochemistry findings, incidence and severity:**
  Rats – Serum protein and albumin concentrations in female rats exposed to 2500 ppm and higher were slightly lower than controls.
  Mice – Blood urea nitrogen concentrations of 5000-ppm females and 10000-ppm males and females were greater than controls.

- **Urinalysis:**
  Rats – At the end of the study, 16 hour urine volumes in 10000 and 20000-ppm male rats were two-fold greater than controls. Urine specific gravity was lower than controls in 5000, 10000 and 20000-ppm males.
  Mice – Decreased urine specific gravity occurred in 10000-ppm females.

- **Mortality and time to death:**
  Rats – no animals died during the study
  Mice – one female (control), 2 males & 1 female (625 ppm), 1 female (1250 ppm), 1 female (2500 ppm), 1 female (5000 ppm), 3 males (10000 ppm) died during the study.

- **Gross pathology incidence and severity:**
  Rats – There were no treatment-related gross lesions.

Mice - There were no treatment-related gross lesions.

- **Organ weight changes:**
  Rats – No biologically significant differences in organ weights were observed
  Mice – The absolute and relative weights of several organs in 5000 and 10000-ppm animals were lower than those in the control group. These findings were attributed to the low body weights in these groups. In males exposed to 5000 or 10000 ppm, weights of the right cauda and right epididymis were significantly lower than those of control males and decreased with increasing exposure level.

- **Histopathology incidence and severity:**
  Rats – A minimal to mild degeneration of the renal papilla was present in 5000, 10000 and 20000-ppm male rats and in one female rat in the 20000 ppm group. Minimal hyperplasia of the transitional epithelium of the urinary bladder was present in 20000-ppm males.
  Mice – There was an exposure and treatment-related increase in the incidence of papillary necrosis. In the cortex of the kidney, there were foci of renal tubule regeneration and fibrosis. These lesions were present in 2500, 5000 and 10000-ppm males and 10000-ppm females. Mild urinary bladder hyperplasia of the transitional epithelium was observed in 5000 and 10000-ppm males and females. In 7/9 female mice from the 10000-ppm group there was also a minimal inflammatory cell infiltration in the urinary bladder mucosa and focal necrosis of the transitional cell epithelium.

Incidence of selected non-neoplastic lesions in rats

	Dose (ppm)					
	0	1250	2500	5000	10000	20000
Male						
Kidney [a]	10	10	10	10	10	10
Degeneration, Papillary [b]	0	0	0	3 (1.0)[c]	6** (1.3)	8** (1.3)
Urinary Bladder	10	10	10	10	10	10
Hyperplasia	0	0	0	0	0	9** (1.0)
Female						
Kidney	10	10	10	10	10	10
Degeneration, Papillary	0	0	0	0	0	1 (1.0)

** Significantly different (P ≤0.01) from the control group by the Fisher exact test. [a] Number of animals with organ examined microscopically. [b] Number of animals with lesion. [c] Average severity grade of lesions in affected rats: 1 = minimal, 2 = mild, 3 = moderate, 4 = marked

Incidence of selected non-neoplastic lesions in mice

	Dose (ppm)					
	0	625	1250	2500	5000	10000
Male						
Kidney [a]	10	10	10	10	10	10
Necrosis, Papillary [b]	0	0	0	5* (1.2)[c]	4* (1.5)	9** (2.2)
Regeneration, Renal Tubule	0	0	0	4* (1.3)	4* (1.5)	7** (2.3)
Fibrosis	0	0	0	4* (1.3)	2 (1.5)	7** (2.1)
Urinary Bladder	10	10	10	10	10	8
Hyperplasia	0	0	0	0	4* (1.0)	7** (2.0)
Female						
Kidney	10	10	10	10	10	10
Necrosis, Papillary	0	0	0	0	0	2 (1.0)
Regeneration, Renal Tubule	0	0	0	0	0	4* (1.8)
Fibrosis	0	0	0	0	0	2 (1.5)
Urinary Bladder	10	10	10	10	10	10
Hyperplasia	0	0	0	0	10** (2.0)	9** (1.6)

* Significantly different (P≤0.05) from the control group by the Fisher exact test. ** P≤0.01. [a] Number of animals with organ examined microscopically. [b] Number of animals with lesion. [c] Average severity grade of lesions in affected mice: 1 = minimal, 2 = mild, 3 = moderate, 4 = marked.

CONCLUSIONS

Remarks:
Rats: Based on lower final mean body weights in the 20000-ppm males and females, the incidences of renal papillary degeneration in 20000-ppm males and females, and hyperplasia of the urinary bladder in 20000-ppm males, the high dose selected for continuous exposure in the 2-year study was 10000 ppm.

Mice: Based on lower final mean body weights and organ weights in 5000 and 10000-ppm males and females and the presence of kidney (papillary necrosis) and urinary bladder lesions in 2500, 5000 and 10000-ppm males and females, the high dose selected for the 2-year study was 1250 ppm.

DATA QUALITY

- 1, Valid without restrictions

Remarks: Well-conducted study performed by American Biogenics Corporation.

REFERENCES:
NTP Technical Report on the Toxicology and Carcinogenesis Studies of 2,2-bis(bromomethyl)-1,3-propanediol (FR-1138, CAS No. 3296-90-0) in F344/N Rats and B6C3F1 Mice (Feed Studies), NTP TR 452, National Toxicology Program, May 1996

OTHER:

- **Last Changed:** October 4 2002
- **Order number for sorting:** 2

Remarks:

HUMAN HEALTH ENDPOINTS
16.3 REPEATED DOSE TOXICITY

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Dow Chemical Company, FR -1138. Purity: approximately 80%, Identity of impurities: monobromoneopentyl triol (6%), tribromoneopentyl alcohol (8%), others (3%)

METHOD

- **Method/guideline followed:** Similar to OECD 453.
- **Test type:** 2 year repeated dose toxicity and carcinogenicity study
- **GLP:** No data
- **Year:** 1979
- **Species:** Rat
- **Strain:** Sprague-Dawley
- **Route of administration:** Oral feed
- **Duration of test:** 2 years
- **Doses/concentration levels:** 0, 5, 100 mg/kg/day
- **Sex:** Male/female
- **Exposure period:** 2 years
- **Frequency of treatment:** Continuous
- **Control group and treatment:** Male/female, 49-50/sex + 5/sex for one year interim kill + 10/sex for tissue analysis
- **Post exposure observation period:** none
- **Statistical methods:** Food consumption, hematology, urinary & clinical chemistry parameters, body & organ weights and organ/body weight ratio data were analyzed by a one-way analysis of variance followed by Dunnett's test (preceded by the sequential outlier's test for food consumption). Data on mortality, palpable masses, gross pathology, histopathology and tumor incidences were analyzed using Fisher's Exact Probability test.

Remarks:

- **Test subjects**
  - **Age at study initiation:** 8-9 weeks

- **No. of animals/sex/dose:** 49-50/sex/dose

- **Study Design**

  - **Vehicle:** Feed

  - **Satellite groups and reasons they were added:** 5/sex/group for 1 year interim kill + 10/sex/group for tissue analysis

  - **Clinical observations performed and frequency:** General health status and possible toxicological response monitored at least weekly during year 1 and almost daily during year 2. Hematological determinations conducted after 90-91 days and 356-357 days of treatment. Additional blood samples collected after 713-714 days and 725 and 731 days (female only). Urine samples collected after 90-91, 356-357 and 713-714 days of treatment. Clinical chemistry parameters were monitored after 94 days, 1 year and 2 years.

  - **Organs examined at necropsy (macroscopic and microscopic):** Year 1 evaluation: Eyes, liver, kidneys, heart, pancreas, spleen, brain, vertebrae with spinal cord, peripheral (sciatic) nerve, pituitary gland, stomach, small intestine, large intestine, cecum, mesenteric lymph node(s), skeletal (thigh) muscle, salivary gland, testes, epididymides, accessory male sex glands, urinary bladder, uretus, ovary trachea, esophagus, aorta, thoracic lymph node(s), thymus, lungs, integument, thyroid gland, parathyroid glands, adipose tissue, adrenal gland(s), sternum, tongue, mandible and any other grossly observed lesion. The weights of the liver, kidneys, brain, heart and testes (males) were also recorded. Year 2 evaluation: As Year 1, plus femoral bone marrow smear from all female rats.

RESULTS

- **NOAEL:** 5 mg/kg/day
- **LOAEL:** 100 mg/kg/day
- **Actual dose received by sex:** The concentration of the test material was adjusted on a weekly basis for the first 3 months, and quarterly thereafter to maintain the designated dose levels on a mg/kg bw/day basis. The results of 8 different analyses conducted on samples collected during months 5, 8, 13, 17, 18, 19, 20 and 22 indicated the following analytical content of test material (expressed as percent of nominal concentrations):

Control	Group	None detected
5 mg/kg/day	Female	95.5±24.7%
5 mg/kg/day	Male	113±46.0%
100 mg/kg/day	Female	97.8±7.2%
100 mg/kg/day	Male	97.8±4.4%

- Toxic response/effects by dose level:

- Statistical results, as appropriate:

Remarks:

- **Body weight:** No toxicologically significant differences in body weight were detected.

- **Food/water consumption:** No consistent deviations in food consumption of males or females at either of the 2 dose levels during the study.

- **Description, severity, time of onset and duration of clinical signs:** None

- **Ophthalmologic findings, incidence and severity:** No data

- **Hematologic findings, incidence and severity:** No toxicologically significant differences between controls and treatment groups.

- **Clinical biochemistry findings, incidence and severity:** No toxicologically significant differences between controls and treatment groups.

- **Mortality and time to death:** Male rats given 5 or 100 mg/kg/day showed no differences in mortality compared to that of the control group. Female rats receiving 100 mg/kg/day had statistically increased mortality rates for months 16 and 17, but this was considered to be of questionable toxicological significance.

- **Gross pathology incidence and severity:** Six of eleven female rats given 100 mg/kg/day showed bilateral diffuse opacity of the lenses. This was not noted in the controls or the group of females given 5 mg/kg/day.

- **Organ weight changes:** Male rats given 100 mg/kg/day killed after 1 year of treatment showed a statistically significant increase in relative liver weights. This was the only toxicologically significant difference between controls and treatment groups.

- **Histopathology incidence and severity:** Statistical increase in the incidence of thyroid retention cyst formation in 100 mg/kg/day male rats, which may or may not have been treatment-related. Female rats given 100 mg/kg/day showed six of eleven rats with bilateral lenticular degeneration of the anterior cortex graded as moderate in degree. In addition, some of the rats showed bilateral degeneration of the posterior cortex and basophilic staining material within this region of the lens. There were several observations noted on examination of the livers of female rats given 100 mg/kg/day that were considered to be related to treatment. These included hepatocellular degenerative changes consisting of increased eosinophilic cytoplasmic homogeneity accompanied by a slight increase in the incidence of livers having several foci or a single area of hepatocellular alteration. The trend towards a slight increased incidence of individual hepatocellular necrosis noted in this group may or may not have been the result of treatment.

  Statistical analysis of the incidence of tumors found at necropsy showed there were no significant differences between control and treatment groups.

CONCLUSIONS

Remarks:

DATA QUALITY

- 2, Valid with restrictions

Remarks: Appears to be a well conducted study, but no GLP information.

REFERENCES:

Keyes, D.G., et al, Results of a two year toxicity and onconogenic study of rats ingesting diets containing dibromoneopentyl glycol (FR -1138), J. Combustion Toxicology, 7, p77, 1980

OTHER:

- **Last Changed:** October 4 2002
- **Order number for sorting:** 3

Remarks:

HUMAN HEALTH ENDPOINTS
17. TOXICITY TO REPRODUCTION

TEST SUBSTANCE

- 2,2-bis(bromomethyl)-1,3-propanediol (CAS No. 3296-90-0)

Remarks: Source: Dow Chemical Company, Lot No.: MM05137-636, Purity: approx 87.3%, Identity of impurities: 2,2-bis (hydroxymethyl)-1-bromo-3-hydroxypropane (6.7%), 2,2-bis (bromomethyl)-1-bromo-3-hydroxypropane (5.5%), 3,3-bis (bromomethyl) oxetane (0.5%), water (0.3±0.1%).

METHOD

- **Method/guideline:** The study was performed using the NTP Fertility Assessment by Continuous Breeding (FACB) system. It consists of four related tasks as follows: Task 1, dose finding; Task 2, continuous breeding phase; Task 3, identification of the affected sex; and Task 4, offspring assessment. Task 1 is conducted to determine suitable doses for the cohabitation study. The test chemical is administered for 14 consecutive days and the maximum tolerated dose (MTD) is estimated. Task 2 is designed to determine the effect of the estimated MTD and two lower dose levels on fertility and reproduction. In this phase, treatment is continued for 18 weeks. Task 3 (a crossover mating trial) is conducted using males and females from the control and high dose groups to determine whether the male, female or both sexes are affected. Task 4 is designed to evaluate reproductive performance in offspring (second generation) from the final Task 2 litters.

- **Type:** Two generation

- **GLP:** Yes

- **Year:** 1986

- **Species:** Mouse

- **Strain:** CD-1 (ICR) BR

- **Route of administration:** Oral feed

- **Doses/concentration levels:** Range Finding Study (Task 1): 0, 0.025, 0.05, 0.1, 0.25, 0.5 % w/w
  Cohabitation Study (Task 2): 0, 0.1, 0.2, 0.4 %w/w
  Crossover Mating Study (Task 3): No chemical treatment during the mating period. 0, 0.4 %w/w during post-mating holding period.
  Second Generation Study (Task 4): 0, 0.1, 0.2, 0.4 %w/w

- **Sex:** Male/female

- **Control group:** Range Finding Study (Task 1): 8 males, 8 females; vehicle control
  Parental (Task 2): 40 males, 40 females; vehicle control
  Crossover Mating Study (Task 3): 20 males, 20 females; vehicle control
  Second Generation Study (Task 4): 20 males, 20 females; vehicle control

- **Frequency of treatment:** Daily, ad libitum

- **Duration of test:** Range Finding Study (Task 1): 2 weeks
  Cohabitation Study (Task 2): 1 week pre-treatment, 14 weeks cohabitation, and 3 weeks holding period
  Crossover Mating Study (Task 3): 1 week cohabitation (maximum), and 3 weeks holding period
  Second Generation Study (Task 4): 74±10 days premating, 7 days (maximum cohabitation), and 3 weeks holding period

- **Premating exposure period for males:** Parental (Task 2): 7 days

F1 (Task 4): 64-84 days

- **Premating exposure period for females:** Parental (Task 2): 7 days
  F1 (Task 4): 64-84 days

- **Statistical methods:** The Cochran-Armitage test was used in Task 2 to assess dose-related trends in fertility. Days between litters were evaluated by Shirley's test. In Task 3 a χ^2 test for homogeneity was used to test for an overall difference in fertility among the cross-mated groups. Pairwise comparisons between control and dosed groups were determined by the Fisher's exact test. Dose group means for the proportion of live pups and sex ratio of pups were tested for overall differences by the Kruskal-Wallis test and for ordered differences by the Jonckheere test. Pairwise comparisons of treatment group means were determined by the Wilcoxon-Mann-Whitney *U* test. Differences in average pup weight were investigated using analysis of covariance (ANCOVA), using avaerage litter size inclusive of live and dead pups as covariant. Pairwise comparisons were performed using a two-tailed *t* test. For organ weights, analysis of covariance was performed to determine least squares treatment group means, which were then tested for overall and pairwise equality by two-sided *F* and *t* tests, respectively. The Kruskal-Wallis and Wilcoxon-Mann-Whitney *U* tests were also employed in the analysis of unadjusted body and organ weights.

Remarks:

- **Test animals**

  - **Parental:** 20 male, 20 female per dose. Age: 11 weeks
  - **F1:** 20 male, 20 female per dose. Age: 64-84 days

- **Test design**

  - **Vehicle:** Substance was administered in feed.

  - **Dosing schedules and pre and post dosing observations periods:**

    Parental (Task 2): Animals fed daily ad libitum for 7 days prior to mating followed by 14 weeks after initial mating and a further 3 week observation period.
    F1 (Task 4): Animals received the substance initially through lactation for 3 weeks until weaning, then fed daily ad libitum until mating at 74±10 days, then for a further 7 days. All animals were necropsied at the end of this period.

- **Mating procedures:** Task 2: 1 male and 1 female per cage. Cohabitation for 14 weeks.
  Task 3: 1 male and 1 female per cage. Cohabitation until vaginal plug was detected or for a maximum of 7 days.
  Task 4: 1 male and 1 female per cage. Cohabitation continued for seven days or until a copulatory plug was found.

- **Standardization of litters:** No

- **Parameters assessed during study P and F1 as appropriate:**

  - **Clinical observations performed and frequency:** Task 2 parental body weights were measured on days –7 (start of dosing), 0 (start of cohabitation), 7, 28, 56, 84 and 112.

    F1 animals (Task 4) were weighed at weaning, the first day of cohabitation and once a week thereafter.

  - **Estrous cycle length and pattern (number of days spent in each phase):** Studied during Task 3 (Parental) and Task 4 (F1).

  - **Sperm examination:** Sperm motility, sperm counts per 'g' caudal tissue, and the incidence of abnormal sperm were examined at the end of Task 3 (parental) and Task 4 (F1)

- **Parameters assessed during study F1 and F2 as appropriate:**

- **Clinical observations performed and frequency:** At least twice per day: Loss of hair, diarrhoea, wounds, hyperactivity, inactivity, tearing or ocular discharge, paralysis, nasal discharge, blood in cage and/or on animal, abnormal posture or death.

- **Others, for example anogenital distance, if performed:** Anogenital distance was used to sex pups

- **Organs examined at necropsy (macroscopic and microscopic):** Female – liver, kidney, lymph nodes, mammary glands, ovary, oviduct, uterus, vagina, urinary bladder and skeletal muscle; Male – kidney, liver, epididymis, prostate gland, seminal vesicles, skin, testes and urinary bladder.

RESULTS

- **F0 NOEL:** 0.1% adults, 0.1% offspring
- **F1 NOEL:** 0.1% adults, 0.1% offspring

- **Actual dose received by dose level by sex:** Male mice in the 0.1, 0.2 and 0.4% dose groups received approximately 141, 274 and 589 mg/kg bw/day. Female dose was not calculated.

- **Parental data and F1 as appropriate (toxic response/effects with NOAEL value). Provide at a minimum qualitative descriptions of elements where dose related observations were seen:** Refer to Remarks

- **Offspring toxicity F1 and F2 as appropriate (toxic response/effects with NOAEL value). Provide, at a minimum, qualitative descriptions of elements where dose related observations were seen:** Refer to Remarks

- **Statistical results, as appropriate:** Refer to Remarks

Remarks:

- **Parental data and F1 as appropriate, provide at a minimum qualitative descriptions of elements where dose related observations were seen:**

- **Body weight:** Parental (Task 2): The average body weights of male and female mice in the 0.4% dose groups were significantly lower ($p<0.05$) than the control groups from week 2 onwards (apart from females at week 14), as shown in the table below.

Treatment Group		Body Weight (g) Group Mean±SE						
		Week 1	Week 2	Week 3	Week 6	Week 10	Week 14	Week 18
Control (0.0 mg/g)	M	33.6±0.30 (40)[a]	35.7±0.32 (40)	34.6±0.29 (40)	35.3±0.30 (40)	36.6±0.35 (40)	37.5±0.38 (40)	38.2±0.44 (39)[b]
	F	26.2±0.22 (40)	28.0±0.24 (40)	30.5±0.24 (40)	34.6±0.39 (40)	49.4±1.01 (40)	40.7±1.16 (40)	45.0±1.02 (40)
0.1% (1.0 mg/g)	M	33.9±0.51 (20)	35.1±0.49 (20)	34.1±0.48 (20)	35.0±0.59 (20)	36.4±0.73 (20)	37.2±0.83 (20)	-[c]
	F	26.0±0.34 (20)	27.1±0.30 (20)	30.2±0.39 (20)	33.7±0.58 (20)	48.1±1.85 (20)	38.6±1.10 (20)	42.0±1.19 (20)
0.2% (2.0 mg/g)	M	34.1±0.37 (20)	35.4±0.39 (20)	34.3±0.38(20)	35.0±0.59 (20)	36.4±0.81 (20)	36.9±0.94 (20)	-[c]
	F	25.8±0.38 (20)	27.1±0.36 (20)	29.3±0.35 (20)[d]	32.8±0.37 (20)[d]	45.7±1.37 (19)[e]	37.6±1.09 (19)	41.0±1.20 (17)[d,f]
0.4% (4.0 mg/g)	M	34.0±0.35 (20)	33.7±0.42 (20)[d]	32.5±0.33 (20)[d]	32.5±0.42 (20)[d]	33.1±0.48 (20)[d]	34.1±0.60 (20)[d]	34.3±0.71 (20)[d]
	F	25.9±0.42 (20)	27.0±0.36 (20)[d]	28.4±0.39 (20)[d]	30.2±0.94 (20)[d]	33.7±1.18 (20)[d]	36.6±1.69 (19)[g]	34.7±1.40 (19)[d]

[a] Number of animals providing the data indicated in parenthesis. [b] One male was sacrificed during week 16 of the study as per instructions of the veterinarian in charge. [c] animals were sacrificed during week 16 of the study. [d] Significantly different ($p<0.05$) from the control group. [e] One female died during week 7 of the study. [f] One female died during week 17 of the study and 1 female was sacrificed during week 17 of the study as per instructions of the veterinarian in charge. [g] One female died during week 11 of the study

Parental (Task 3): The body weights of both males and females from the 0.4% groups were significantly lower than the animals in the control group throughout the 4-week study, as shown in the table below:

Treatment Group		Body Weight (g) Group Mean±SE			
		Week 22	Week 23	Week 24	Week 25
Control (0.0 mg/g)	M	40.9±0.49 (39)[a]	38.7±0.47 (39)	39.3±0.48 (39)	38.7±0.51 (39)
	F	37.6±0.43 (40)	37.7±0.50 (40)	43.2±0.81 (40)	46.4±1.65 (40)
0.4% (4.0 mg/g)	M	35.3±0.88 (20)[b]	34.4±0.72 (20)[b]	34.1±0.83 (20)[b]	33.8±0.90 (20)[b]
	F	31.0±0.60 (18)[b]	31.7±0.61 (18)[b]	33.7±0.81 (18)[b]	38.4±1.61 (18)[b]

[a] Number of animals providing the data indicated in parenthesis. [b] Significantly different ($p<0.05$) from the control group

F1 (Task 4): Pups in the 3 dose groups were smaller than the control group at weaning as well as through maturation. As shown in the table below, this difference was significant for both males and females in the 0.4% dose groups at all time intervals.

Treatment Group		Body Weight (g) Group Mean±SE				
		Week 19-21[a]	Week 28	Week 29	Week 30	Week 31
Control (0.0 mg/g)	M	11.9±0.49 (20)[b]	33.9±0.59 (20)	34.0±0.60 (20)	35.4±0.64 (20)	36.4±0.66 (20)
	F	11.4±0.44 (40)	26.6±0.56 (19)[c]	30.0±0.64 (20)	37.5±0.82 (20)	43.7±2.85 (20)
0.1% (1.0 mg/g)	M	10.2±0.40 (20)[d]	32.8±0.62 (20)	33.3±0.52 (20)	33.6±0.55 (20)[d]	34.6±0.56 (20)[d]
	F	9.9±0.39 (20)[d]	24.5±0.41 (20)[d]	28.5±0.40 (20)	34.8±0.64 (20)[d]	41.6±2.58 (20)
0.2% (2.0 mg/g)	M	10.5±0.45 (20)[d]	31.5±0.62 (20)[d]	31.7±0.59(20)[d]	32.4±0.61 (20)[d]	33.1±0.60 (20)[d]
	F	10.2±0.43 (20)	24.5±0.50 (20)[d]	28.3±0.58 (20)	33.8±0.91 (20)[d]	40.7±2.67 (19)
0.4% (4.0 mg/g)	M	9.0±0.53 (20)[d]	25.5±0.59 (20)[d]	26.2±0.56 (20)[d]	26.3±0.67 (20)[d]	26.9±0.80 (20)[d]
	F	8.8±0.47 (20)[d]	21.7±0.37 (20)[d]	24.3±0.39 (20)[d]	28.5±0.70 (20)[d]	32.1±1.77 (20)[d]

[a] Represents average pup weight at weaning; pups were weaned during weeks 19 to 21 of the study. [b] Number of animals providing the data indicated in parenthesis. [c] Due to technical error, one female mouse was not weighed during week 28. [d] Significantly different ($p<0.05$) from the control group.

- **Food/water consumption:** There were no statistically significant differences in the average daily consumption of control/dose feed by animals in the control and treatment groups (parental or F1) in this study.

- **Description, severity, time of onset and duration of clinical signs:** None

- **Fertility index:** There was no significant difference in the fertility index values between control and dose group animals during Task 2 (Parental) or between control and dose group animals during Task 4 (F1). The fertility index of the control male x 0.4% dose females was significantly different ($p<0.01$) from the control male x control female group in Task 3 (Parental). The data are presented in the tables below.

Fertility of Pairs during Continuous Breeding (Parental, Task 2)

Treatment group	No. Fertile/No. Cohabited	Fertility Index (%)[a]
Control	39/40	97
0.1%	19/20	95
0.2%	19/19[b]	100
0.4%	19/19[c]	100

[a] Fertility Index (%) = 100 x No. Fertile/No. Cohabited. [b] One female died during week 7 of the study; data were excluded. [c] One female died during week 11 of the study; data were excluded

Mating and fertility of Pairs after a Mating trial to determine the Affected Sex (Parental, Task 3)

Treatment group	No. with Copulatory Plugs/ No. Cohabited	Mating Index (%)[a]	No. Fertile/ No. Cohabited	Fertility Index (%)[b]
Control Male x Control Female	19/20[c]	95	17/20	85
Control Male x 0.4% Female	17/18	94	6/18	33[d]
0.4% Male x Control Female	17/20	85	16/20	80

[a] Mating Index (%) = 100 x No. with Copulatory Plugs/No. Cohabited. [b] Fertility Index (%) = 100 x No. Fertile/No. Cohabited. [c] Although not detected by direct means, 2 females were scored copulatory plug-positive based on delivery of litters. [d] Significantly different ($p<0.01$) from the control male x control female group.

Reproductive Performance (F1, Task 4)

Treatment	No. with Copulatory Plugs/	Mating Index	No. Fertile/	Fertility Index

group	No. Cohabited	(%)[a]	No. Cohabited	(%)[b]
Control	20/20	100	19/20	95
0.1%	20/20[c]	100	18/20	90
0.2%	20/20	100	18/20	90
0.4%	20/20	100	18/20	90

[a] Mating Index (%) = 100 x No. with Copulatory Plugs/No. Cohabited. [b] Fertility Index (%) = 100 x No. Fertile/No. Cohabited
[c] Although not detected by direct means, 2 females were scored copulatory plug-positive based on delivery of litters.

- **Precoital interval (with number of days until mating and number of estrous periods until mating):** No data

- **Duration of gestation (calculated from day 0 of pregnancy):** The number of days to litter for the 0.4% dose group was significantly increased over control group values for the first four litters. It was also noted that there was a decrease in dams delivering fourth and fifth litters in the 0.4% treatment group.

Days to Litter (Task 2)

Treatment Group	Days to litter				
	Litter 1	Litter 2	Litter 3	Litter 4	Litter 5
Control	20.62±0.17 (39)[a]	20.13±0.23 (39)	20.28±0.14 (39)	20.13±0.08 (39)	20.58±0.37 (38)
0.1%	20.47±0.32 (19)	20.32±0.61 (19)	20.42±0.27 (19)	20.11±0.13 (10)	20.00±0.08 (19)
0.2%	21.30±0.46 (20)	19.85±0.13 (20)	20.32±0.11 (19)	20.84±0.41 (19)	20.67±0.44 (18)
0.4%	26.70±2.40 (20)*	27.80±3.35 (20)*	22.74±1.45 (19)*	22.06±1.21 (16)[b],*	20.73±0.19 (11)

[a] All data are expressed as means±SE, with the number of dams producing litters in parenthesis. [b] One female died during Week 11 of the study. * Denotes significantly different from the control ($p<0.05$)

- **Gestation index (live litters/pregnancies):** No data

- **Change in lactation:** No data

- **Changes in estrous cycles:** A detailed vaginal cytology evaluation was undertaken at the end of Task 3 (Parental). The average estrous cycle length in the treated animals was 4.9 days vs. a control value of 4.7 days. Treatment with the test substance had no apparent effect on estrual cyclicity.

Summary of Data from Vaginal Cytology Studies (Parental, Task 3)

Treatment group	Terminal Body Weight (g)	Relative Frequency of Estrous Stages					Estrous Cycle Length (days)
	Group Mean±SE	%P	%E	%M	%D	%NC	Group Mean±SE
Control	38.0±0.56 (38)[a,b]	13.0 (39)	20.0 (39)	23.0 (39)	44.1 (39)[c]	0.0 (39)	4.69±0.12 (26)[b,d]
0.4%	31.4±0.62 (17)[e]	18.5 (17)	22.7 (17)	22.7 (17)	36.1 (17)[f]	0.0 (17)	4.93±0.20 (14)[g]

[a] Number of animals providing the data indicated in parenthesis. [b] In the control group, 1 mouse was found dead on day 5 of vaginal smearing. Data from 4 days of vaginal smears are included. [c] For 7 consecutive days, 3 animals in the control group remained in diestrus stage. [d] In 12 out of the 39 animals in the control group, estrous cycle length was 7 days or not clear. [e] Significantly different ($p<0.01$) from the control group. [f] For 7 consecutive days, 2 animals in the 0.4% dose group remained in diestrus stage. [g] In 3 out of the 17 animals in the 0.4% dose group, estrous cycle length was not clear. P = Proestrus; E = Estrus; M = Metestrus; D = Diestrus; NC = Not Clear

In the F1 study (Task 4) vaginal smears were prepared for 7 days prior to necropsy. The average estrous cycle varied between 5.0 to 5.2 days for animals in the control and 3 dose groups. Estrual cyclicity was not affected at the 0.1 and 0.2% levels but at the 0.4% level relative frequency of the diestrus phase was 43% vs. a control value of 25%.

Summary of Data from Vaginal Cytology Studies (F1, Task 4)

Treatment group	Terminal Body Weight (g)	Relative Frequency of Estrous Stages					Estrous Cycle Length (days)
	Group Mean±SE	%P	%E	%M	%D	%NC	Group Mean±SE
Control	32.2±0.71 (20)[a]	22.1 (20)	32.1 (20)	20.7 (20)	25.0 (20)	0.0 (20)	5.18±0.15 (17)[b]
0.1%	30.2±0.35 (20)[c]	14.3 (20)	34.3 (20)	22.9 (20)	28.6 (20)	0.0 (20)	4.95±0.12 (19)[d]
0.2%	30.4±0.56 (20)	20.0 (20)	35.0 (20)	17.1 (20)	27.9 (20)	0.0 (20)	5.21±0.14 (19)[e]
0.4%	26.5±0.41 (20)[f]	16.4 (20)	25.0 (20)	15.7 (20)	42.9 (20)	0.0 (20)	4.94±0.15 (18)[g]

a Number of animals providing the data indicated in parenthesis. [b] In 3 out of the 20 animals in the control group, estrous cycle length was 7 days. [c] Significantly different ($p<0.05$) from the control group. [d] In 1 out of the 20 animals in the 0.1% dose group, estrous cycle length was 7 days. [e] In 1 out of the 20 animals in the 0.2% dose group, estrous cycle length was not clear. [f] Significantly different ($p<0.01$) from the control group. [g] In 2 out of the 20 animals in the 0.4% dose group, estrous cycle length was 7 days or not clear. P = Proestrus; E = Estrus; M = Metestrus; D = Diestrus; NC = Not Clear

- **Effects on sperm:** Sperm assessment studies following Task 3 (Parental) showed that treatment with the test substance had no significant effect ($p>0.05$) on sperm motility, sperm counts per 'g' caudal tissue, or the incidence of abnormal sperm (see below).

Summary of Data from Sperm Studies (Parental, Task 3)

Treatment group	Weight				Sperm Motility (%)	Sperm Density x 10^{6} [a]	Abnormal Sperm (%)
	Body (g)	R. Caudal (mg)	R. Epididymal (mg)	R. Testicular (g)			
Control	40.65 ±0.52 (39)[b,c]	19.564±0.300 (39)	53.515±0.737 (39)	0.140±0.004 (39)	94.05±0.40 (39)	1102±46 (39)	3.07±0.36 (39)
0.4%	35.67±1.01 (19)[d,f]	17.705±0.645 (19)[e]	50.500±1.217 (19)	0.138±0.004 (19)	94.12±0.80 (19)	1076±42 (19)	2.92±0.24 (19)

[a] Per g caudal tissue. [b] Number of animals providing the data indicated in parenthesis. [c] Mean±SE. [d] Significantly different ($p<0.01$) from the control group. [e] Significantly different ($p<0.05$) from the control group. [f] Severe internal infection noted at necropsy for 1 mouse: all data were excluded

Sperm assessment studies following Task 4 (F1) showed that treatment with the test substance at up to 0.4% dose level has no significant effect ($p>0.05$) on sperm motility or the incidence of abnormal sperm in the second generation CD-1 mice. The sperm density was reduced by approximately 14% in the 0.4% dose group (see below).

Summary of Data from Sperm Studies (F1, Task 4)

Treatment group	Weight				Sperm Motility (%)	Sperm Density x 10^{6} [a]	Abnormal Sperm (%)
	Body (g)	R. Caudal (mg)	R. Epididymal (mg)	R. Testicular (g)			
Control	37.57±0.71 (20)[b,c]	18.850±0.696 (20)	49.040±1.518 (20)	0.140±0.004 (20)	88.6±2.95 (20)	1307±66 (20)	3.31±0.49 (20)
0.1%	35.73±0.64 (20)	18.820±0.411 (20)	49.740±1.481 (20)	0.139±0.005 (20)	93.1±0.90 (20)	1427±49 (20)	3.28±0.43 (20)
0.2%	34.37±0.61 (20)[d]	18.550±0.594 (20)	47.800±1.107 (20)	0.139±0.005 (20)	92.3±0.98 (20)	1400±91 (20)	2.56±0.43 (20)
0.4%	28.24±0.87 (20)[d]	14.530±0.622 (20)[d]	39.230±1.413 (20)[d]	0.117±0.004 (20)[d]	91.9±0.91 (20)	1124±76 (20)[e]	3.70±0.59 (20)

[a] Per g caudal tissue. [b] Number of animals providing the data indicated in parenthesis. [c] Mean±SE. [d] Significantly different ($p<0.01$) from the control group. [e] Significantly different ($p<0.05$) from the control group

- **Hematological findings incidence and severity:** No data
- **Clinical biochemistry findings incidence and severity:** No data
- **Mortality:** Task 2: Two females (0.2% dose group) died during weeks 7 & 17 and other female from this group was sacrificed during week 17. One female (0.4% dose group) died during week 11.
- **Gross pathology incidence and severity:** No data
- **Clinical biochemistry findings incidence and severity:** No data
- **Number of implantations:** No data
- **Number of corpora lutea (recommended):** No data

- **Ovarian primordial follicle counts:** No data

- **Organ weight changes:** For parental animals necropsied at the end of Task 3, the group mean whole body, liver and kidney weights of female mice in the 0.4% dose group were significantly lower ($p<0.05$) than the corresponding control values. The reduction was no longer apparent when organ weights were adjusted for body weight at necropsy. For males in the 0.4% dose group, whole body, kidney, seminal vesicles and right caudal weights were significantly lower ($p<0.05$) than the corresponding control values. These differences were not apparent when organ weights were adjusted for body weight at necropsy.

- **Histopathology incidence and severity:** For parental animals necropsied at the end of Task 3, significant gross lesions were seen in the kidneys of males and females treated with the substance. Pathologic examination revealed significantly increased incidence of glomerular atrophy, degeneration, dilation, and regeneration of tubules, renal papillary necrosis, interstitial nephritis, and mineralisation in animals treated with the substance.

  A detailed histopathologic examination of F1 mice at the end of Task 4 revealed significant treatment-related gross lesions in the female mice. Changes were also observed in the kidney of many on the male animals from the 0.2 and 0.4% dose groups.

- **Offspring toxicity F1 and F2, as appropriate, provide as a minimum qualitative descriptions of elements where dose related observations were seen:**

- **Litter size and weights, sex and sex ratios:** In the continuous breeding study (Task 2) there was a significant decrease ($p<0.01$) in the mean number of litters per fertile pair at the 0.4% dose level. Other reproductive parameters affected ($p<0.01$) at this dose level were the total number of live pups per litter, the proportion of live pups born alive and the adjusted live pup weight. At the 0.2% dose level, both absolute (except for female pups) and adjusted live pup weights were significantly decreased ($p<0.05$).

Reproductive Performance of Fertile Pairs During Continuous Breeding (Parental, Task 2)

Reproductive Parameter [a]	Treatment Group			
	Control	0.1%	0.2%	0.4%
Litters per Pair	4.97±0.03 (39)[b]	5.00±0.00 (19)	4.95±0.05 (19)	4.37±0.21 (19)[c]
Live Pups per Litter				
Male	6.12±0.23 (39)	6.24±0.35 (19)	5.64±0.32 (19)	3.47±0.27 (19)[c]
Female	6.11±0.19 (39)	6.19±0.30 (19)	5.68±0.32 (19)	3.34±0.30 (19)[c]
Combined	12.23±0.37 (39)	12.43±0.49 (19)	11.32±0.59 (19)	6.81±0.49 (19)[c]
Proportion of Pups Born Alive	0.99±0.01 (39)	0.99±0.01 (19)	0.99±0.00 (19)	0.97±0.01 (19)[c]
Sex of Pups Born Alive (Male/Total)	0.50±0.01 (39)	0.50±0.02 (19)	0.50±0.01 (19)	0.51±0.03 (19)
Live Pup Weight (g)				
Male	1.61±0.02 (39)	1.57±0.02 (19)	1.53±0.02 (19)[c]	1.56±0.02 (19)
Female	1.53±0.01 (39)	1.52±0.02 (19)	1.49±0.02 (19)	1.53±0.02 (19)
Combined	1.57±0.01 (39)	1.55±0.02 (19)	1.51±0.02 (19)[d]	1.54±0.02 (19)
Adjusted Live Pup Weight (g)[e]				
Male	1.63±0.01 (39)	1.60±0.02 (19)	1.53±0.02 (19)[c]	1.48±0.02 (19)[c]
Female	1.56±0.01 (39)	1.55±0.02 (19)	1.49±0.02 (19)[c]	1.45±0.02 (19)[c]
Combined	1.59±0.01 (39)	1.57±0.02 (19)	1.51±0.02 (19)[c]	1.47±0.02 (19)[c]

[a] Mean±SE. [b] Number of fertile pairs providing the data indicated in parenthesis. [c] Significantly different ($p<0.01$) from the control group. [d] Significantly different ($p<0.05$) from the control group. [e] Least squares estimate of mean pup weight adjusted for average litter size±SE

In Task 3 (crossover study), there was a significant difference ($p<0.05$) in the number of live pups between the control male x 0.4% dose female and the 0.4% dose male x control female group. Adjusted pup weights in the former group were also significantly reduced ($p<0.05$) relative to the control male x control female group.

Reproductive Performance of Fertile Pairs after a Mating Trial to Determine Affected Sex (Parental, Task 3)

Reproductive Parameter [a]	Treatment Group		
	Control Male x Control Female	Control Male x 0.4% Female	0.4% Male x Control Female

Live Pups per Litter			
Male	5.00±0.39 (17)[b]	3.17±0.75 (06)[c,d]	5.69±0.42 (16)
Female	4.71±0.80 (17)	3.67±0.84 (06)	5.06±0.49 (16)
Combined	9.71±1.01 (17)	6.83±1.40 (06)[d]	10.75±0.63 (16)
Proportion of Pups Born Alive	0.96±0.03 (17)	0.91±0.07 (06)	0.98±0.01 (16)
Sex of Pups Born Alive (Male/Total)	0.53±0.05 (17)	0.47±0.07 (06)	0.54±0.03 (16)
Live Pup Weight (g)			
Male	1.68±0.04 (16)[e]	1.69±0.08 (06)	1.62±0.04 (16)
Female	1.60±0.05 (16)[f]	1.58±0.05 (06)	1.55±0.03 (16)
Combined	1.66±0.04 (17)	1.63±0.07 (06)	1.59±0.03 (16)
Adjusted Live Pup Weight (g)[e]			
Male	1.70±0.03 (16)	1.57±0.05 (06)[c]	1.65±0.03 (16)
Female	1.61±0.03 (16)	1.48±0.05 (06)[c]	1.59±0.03 (16)
Combined	1.66±0.02 (17)	1.53±0.04 (06)[c]	1.63±0.03 (16)

[a] Mean±SE. [b:] Number of fertile pairs providing the data indicated in parenthesis. [c] Significantly different ($p<0.05$) from the control male x control female group. [d] Significantly different ($p<0.05$) from the 0.4% male x control female group[e] One litter had no live male pups. [f] One litter had no live female pups. [g] Least squares estimate of mean pup weight adjusted for average litter size±SE

In the F1 reproduction study (Task 4) a dose-related decrease was noted with respect to the average number of live pups per litter. The proportion of pups born alive and average pup weight was not affected by treatment with the substance ($p>0.05$). The adjusted pup weight values were significantly decreased ($p<0.05$) at the highest dose level.

Reproductive Performance of Second Generation Fertile Pairs (F1, Task 4)

Reproductive Parameter [a]	Treatment Group			
	Control Male x Control Female	0.1% Male x 0.1% Female	0.2% Male x 0.2% Female	0.4% Male x 0.4% Female
Live Pups per Litter				
Male	6.42±0.55 (19)[b]	5.44±0.34 (18)[c]	5.89±0.60 (18)	3.89±0.51 (18)[d]
Female	5.58±0.47 (19)	6.22±0.32 (18)	4.44±0.70 (18)	4.17±0.54 (18)
Combined	12.00±0.50 (19)	11.67±0.43 (18)	10.33±1.09 (18)	8.06±0.65 (18)[d]
Proportion of Pups Born Alive	1.00±0.00 (19)	1.00±0.00 (18)	0.94±0.06 (18)	0.99±0.01 (18)
Sex of Pups Born Alive (Male/Total)	0.53±0.04 (19)	0.46±0.02 (18)	0.61±0.04 (17)[e]	0.47±0.05 (18)
Live Pup Weight (g)				
Male	1.57±0.02 (19)	1.57±0.03 (18)	1.56±0.05 (17)	1.48±0.03 (17)[c,f]
Female	1.52±0.03 (19)	1.52±0.02 (18)	1.51±0.05 (16)[g]	1.50±0.05 (18)
Combined	1.55±0.02 (19)	1.54±0.02 (18)	1.55±0.05 (17)	1.50±0.04 (18)
Adjusted Live Pup Weight (g)[e]				
Male	1.61±0.03 (19)	1.60±0.03 (18)	1.57±0.03 (17)	1.40±0.03 (17)[d]
Female	1.57±0.03 (19)	1.55±0.03 (18)	1.53±0.03 (16)[g]	1.39±0.03 (18)[d]
Combined	1.60±0.02 (19)	1.58±0.02 (18)	1.55±0.03 (17)	1.40±0.03 (18)[d]

[a] Mean±SE. [b] Number of fertile pairs providing the data indicated in parenthesis. [c] Significantly different ($p<0.05$) from the control group. [d] Significantly different ($p<0.01$) from the control group. [e] One litter contained no live pups. [f] One litter contained no live male pups. [g] One litter contained no live female pups. [h] Least squares estimate of mean pup weight adjusted for average litter size±SE

- **Viability index (pups surviving 4 days/total births) and survival at 14 days:** The final litter (Task 2) from the control and all 3 dose groups was kept to check the effect of the substance on pup survival. Treatment had no apparent effect on pup survival, although it should be stressed that the average litter sizes in the 0.2 and 0.4% dose groups were significantly smaller ($p<0.05$) apparently due to the toxicity of the test substance. Viability index was not calculated in the study report.

Summary of Pup Survival (Task 2, Final litter)

Parameter [a]		Treatment Group			
		Control	0.1%	0.2%	0.4%
Number of Breeding Pairs		40	20	19	19
Number of Litters Born		38	19	18	19
Total Live Pups per Litter					
Age (days)	0	12.29±0.51 (38)[b]	13.32±0.64 (19)	11.00±0.64 (18)	5.74±0.88 (19)[c]
	4	11.08±0.59 (38)	11.32±1.19 (19)	10.33±0.67 (18)	5.00±0.95 (19)[c]
	14	10.66±0.67 (38)	10.95±1.15 (19)	10.41±0.67 (17)[d]	4.95±0.93 (19)[c]
Live Male Pups per Litter					
Age (days)	0	6.03±0.44 (38)	6.42±0.51 (19)	5.22±0.53 (18)	3.16±0.54 (19)[c]
	4	5.32±0.42 (38)	5.47±0.68 (19)	4.83±0.50 (18)	2.84±0.59 (19)[c]
	14	5.16±0.46 (38)	5.21±0.63 (19)	4.94±0.52 (17)[d]	2.84±0.59 (19)[e]
Live Female Pups per Litter					
Age (days)	0	6.26±0.40 (38)	6.89±0.51 (19)	5.78±0.47 (18)	2.58±0.50 (19)[c]
	4	5.76±0.42 (38)	5.84±0.78 (19)	5.50±0.45 (18)	2.16±0.52 (19)[c]
	14	5.50±0.44 (38)	5.74±0.75 (19)	5.47±0.46 (17)[d]	2.11±0.49 (19)[e]

[a] Mean±SE. [b] Number of fertile pairs providing the data indicated in parenthesis. [c] Significantly different ($p<0.01$) from the control group. [d] One litter, along with the dam, was sacrificed on postnatal day 9 as per instructions of veterinarian in charge. [e] Significantly different ($p<0.05$) from the control group

- **Effects on offspring (grossly visible abnormalities):** None
- **Postnatal growth, growth rate:** No data
- **Vaginal opening (F) or preputial separation (M):** No data
- **Other observations, for instance anogenital distance, if measured:** No data
- **Organ weights:** No data for F2
  - **Gross pathology:** No data for F2

CONCLUSIONS

Remarks: The substance administered in feed at the 0.4% dose level adversely affected reproduction in CD-1 mice. Continued treatment at this dose level also resulted in a significant drop in body weight. The crossover mating study showed that it was the reproductive performance of female mice that was affected by treatment with the substance at the 0.4% dose level. The number of live pups per litter delivered by the 0.4% female x control male group being significantly lower than the 0.4% male x control female group. The reproductive performance of second generation CD-1 mice exposed to the substance at the 0.4% dose level was adversely affected with respect to the number of live pups per litter and the adjusted live pup weight. Second generation animals in the 0.4% group weighed less than the control group at weaning, through maturation and at necropsy.

DATA QUALITY

- **Reliabilities:** 1, Valid without restrictions

Remarks: Well-conducted study performed by Environmental Health Research and Testing, Inc.

REFERENCES (Free Text)

Gulati, D.K., et al, Environmental Health Research & Testing, Inc., 2,2-bis(bromomethyl)-1,3-propanediol: Reproduction and fertility assessment in CD-1 mice when administered in feed, NTP Contract # N01-ES-2-5013, Report No. NTP-86-063, January 1986

Treinen, K.A., et al, Reproductive toxicity of 2,2-bis(bromomethyl)-1,3-propanediol in a continuous breeding protocol in Swiss (CD-1) mice, Fundamental & Applied Toxicology, 13, 245-255, 1989

OTHER

- Last Changed: October 4 2002
- Order number for sorting: 1

Remarks: